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(54) Title: NOVEL NEUROTROPHIC FACTORS

(57) Abstract: The invention relates to neublastin neurotrophic factor polypeptides, nucleic acids encoding neublastin polypeptides, and antibodies that bind specifically to neublastin polypeptides, as well as methods of making and methods of using the same.

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## NOVEL NEUROTROPHIC FACTORS

### FIELD OF THE INVENTION

The invention relates to neurotrophic factor polypeptides, nucleic acids encoding neurotrophic factor polypeptides, and antibodies that bind specifically to neurotrophic factors.

5

### BACKGROUND OF THE INVENTION

Neurotrophic factors are naturally-occurring proteins which promote survival, maintain phenotypic differentiation, prevent degeneration, and enhance the activity of neuronal cells and tissues. Neurotrophic factors are isolated from neural tissue and from non-neural tissue that is

10 innervated by the nervous system, and have been classified into functionally and structurally related groups, also referred to as families, superfamilies, or subfamilies. Among the neurotrophic factor superfamilies are the fibroblast growth factor, neurotrophin, and transforming growth factor- $\beta$  (TGF- $\beta$ ) superfamilies. Individual species of neurotrophic factors are distinguished by their physical structure, their interaction with their composite

15 receptors, and their affects on various types of nerve cells. Classified within the TGF- $\beta$  superfamily (Massague, *et al.*, 1994, *Trends in Cell Biology*, 4:172-178) are the glial cell line-derived neurotrophic factor ligands ("GDNF"; WO 93/06116, incorporated herein by reference), which include GDNF, persephin ("PSP"; Milbrandt *et al.*, 1998, *Neuron* 20:245-253, incorporated herein by reference) and neurturin ("NTN"; WO 97/08196, incorporated

20 herein by reference). The ligands of the GDNF subfamily have in common their ability to induce signalling through the RET receptor tyrosine kinase. These three ligands of the GDNF subfamily differ in their relative affinities for a family of neurotrophic receptors, the GFR $\alpha$  receptors.

Due to the effects of neurotrophic factors on neuronal tissue, there remains a need to

25 identify and characterise additional neurotrophic factors for diagnosing and treating disorders of the nervous system.

### SUMMARY OF THE INVENTION

This invention relates to a novel neurotrophic factor herein called "neublastin," or "NBN." Neublastin is classified within the GDNF subfamily because it shares regions of

30 homology with other GDNF ligands (see Tables 3 and 4, *infra*) and because of its ability to

interact with RET (see, *e.g.*, Airaksinen et al., Mol. Cell. Neuroscience, 13, pp. 313-325 (1999)), neublastin is a novel and unique neurotrophic factor. Unlike other GDNF ligands, neublastin exhibits high affinity for the GFR $\alpha$ 3-RET receptor complex and has unique subregions in its amino acid sequence.

5           Included in the invention are NBN nucleic acids and their encoded polypeptides. NBN polypeptides are biologically active as dimers, and have little to any activity as monomers. In one aspect, the invention provides a truncated neublastin polypeptide, wherein the amino terminus of the truncated neublastin polypeptide lacks one or more amino-terminal amino acids of a mature neublastin polypeptide. Preferably, the truncated neublastin polypeptide, when  
10           dimerized, binds to a RET polypeptide. Preferably, the truncated active neublastin polypeptide induces dimerization of the RET polypeptide.

          In some embodiments, the truncated neublastin polypeptide includes seven cysteine residues at positions corresponding to positions 16, 43, 47, 80, 81, 109, and 111 of the neublastin polypeptide of SEQ ID NO:12 (mature 113AA), which correspond, *e.g.*, to positions  
15           43, 70, 74, 107, 108, 136, 138 of SEQ ID NO:9. SEQ ID NO:9 is numbered such that residues -80 through -1 correspond to the prepro section of the translated NBN polypeptide, and residues 1-140 correspond to the 140 amino acids of mature NBN140. This numbering scheme is presented as a matter of convenience only, and no inference is to be drawn from it with regards to any one or more forms of prepro, pro, mature or truncated NBN disclosed herein.

20           Also within the invention is a polypeptide that includes the amino acid sequence of a truncated neublastin polypeptide. The amino acid sequence of the truncated neublastin polypeptide is less than 113 amino acids in length and includes an amino acid sequence at least 70% homologous to amino acids 28-140 of SEQ ID NO:9.

          In some embodiments, the amino acid sequence of the truncated neublastin polypeptide  
25           is at least 80% homologous to amino acids 42-140 of SEQ ID NO:9. More preferably, the amino acid sequence of the truncated neublastin polypeptide is at least 90% homologous to amino acids 42-140 of SEQ ID NO:9. Even more preferably, the amino acid sequence of the neublastin polypeptide is at least 95% homologous to amino acids 42-140 of SEQ ID NO:9. In a further embodiment, the amino acid sequence of the neublastin polypeptide is at least 99%  
30           homologous to amino acids 42-140 of SEQ ID NO:9. In most preferred embodiments, the amino acid sequence of the truncated neublastin polypeptide comprises amino acids 42-140 of

SEQ ID NO:9. In some embodiments, the amino acid sequence of the truncated neublastin polypeptide consists essentially of 99 amino acids of SEQ ID NO:48.

In some embodiments, the amino acid sequence of the truncated neublastin polypeptide is at least 80% homologous to amino acids 39-140 of SEQ ID NO:9. Preferably, the amino acid sequence of the neublastin polypeptide is at least 90% homologous to amino acids 39-140 of SEQ ID NO:9. More preferably, the amino acid sequence of the neublastin polypeptide is at least 95% homologous to amino acids 39-140 of SEQ ID NO:9. In most preferred embodiments, the amino acid sequence of the truncated neublastin polypeptide comprises amino acids 39-140 of SEQ ID NO:9. In a further embodiment, the amino acid sequence of the neublastin polypeptide is at least 99% to amino acids 39-140 of SEQ ID NO:9. In some embodiments, the amino acid sequence of the truncated neublastin polypeptide consists essentially of 102 amino acids of SEQ ID NO:45.

In further embodiments, the amino acid sequence of the truncated neublastin polypeptide is identical to, or at least 80%, 90%, 95% or 99% homologous to, amino acids 29-140 of SEQ ID NO:9, and consists essentially of 112 amino acids. In alternative embodiments, the above mentioned identity or percent homology is to amino acids 30-140 of SEQ ID NO:9, 31-140 of SEQ ID NO:9, 32-140 of SEQ ID NO:9, 33-140 of SEQ ID NO:9, 34-140 of SEQ ID NO:9, 35-140 of SEQ ID NO:9, 36-140 of SEQ ID NO:9, 37-140 of SEQ ID NO:9, 38-140 of SEQ ID NO:9, 40-140 of SEQ ID NO:9 or 41-140 of SEQ ID NO:9. In these embodiments, the amino acid sequence of the truncated neublastin polypeptide consists essentially of 111, 110, 109, 108, 107, 106, 105, 104, 103, 101 or 100 amino acids, respectively. In specific embodiments, the truncated neublastin polypeptide is the polypeptide of any one of SEQ ID NOS:35, 36, 37, 38, 39, 40, 41, 42, 43, 44, 45, 46, 47 or 48.

The truncated neublastin polypeptide can be obtained by providing a mature neublastin polypeptide such as NBN113, and contacting the mature neublastin polypeptide with at least one protease under conditions sufficient to produce the truncated neublastin polypeptide. Preferably, the truncated neublastin polypeptide is produced as an exoprotease neublastin polypeptide digestion product by contacting the mature neublastin polypeptide with at least one exoprotease. A preferred protease is any one of aminopeptidase, Endo Lys C, and trypsin. In some embodiments, the method includes further contacting the exopeptidase neublastin polypeptide digestion product with a dipeptidyl peptidase.

In one embodiment, the truncated neublastin polypeptide is a glycosylated polypeptide. In an alternative embodiment, the truncated neublastin polypeptide is not glycosylated.

Also within the invention is a nucleic acid that includes a polypeptide that includes the amino acid sequence of a truncated Neublastin polypeptide. In some embodiments, the nucleic acid hybridizes specifically under high stringency solution hybridization conditions to a nucleic acid encoding a variant neublastin polypeptide.

The nucleic acid encoding a truncated neublastin polypeptide can be used by introducing the nucleic acid into a cell and causing a polypeptide encoded by the nucleic acid to be expressed in a cell. If desired, the method can include the step of administering the nucleic acid to an animal, and causing the polypeptide to be expressed in the animal.

The nucleic acid encoding a truncated neublastin polypeptide can be provided as a vector, *e.g.*, an expression vector. The vector can be used to express the encoded truncated neublastin polypeptide.

The invention also includes a cell transformed with a nucleic acid encoding a polypeptide that includes a truncated neublastin polypeptide. The cell can be, *e.g.* a mammalian cell, a fungal cell, a yeast cell, an insect cell, and a bacterial cell. A preferred mammalian cell is a Chinese hamster ovary cell, or a cell derived from the mammalian central nervous system.

In a further aspect, the invention includes a method of making a truncated neublastin polypeptide by expressing a nucleic acid encoding a truncated neublastin polypeptide. Preferably, the method includes the step of culturing a cell comprising the nucleic acid in a culture medium which permits the production of the truncated neublastin polypeptide. The method can also include the step of recovering the polypeptide from the culture medium. Also provided by the invention is a truncated neublastin polypeptide, (*e.g.*, a purified protein) obtained by the method.

Also provided by the invention is a pharmaceutical composition that includes a truncated neublastin polypeptide and a pharmaceutically acceptable carrier. Also within the invention is a pharmaceutical composition comprising a nucleic acid encoding a truncated neublastin polypeptide and a pharmaceutically acceptable carrier.

In a still further aspect, the invention provides a method of administering the truncated neublastin polypeptide by delivering the polypeptide to an isolated cell or *in vivo* to a mammal (such as a human). Preferably, the administration *in vivo* comprises systemic administration.

The mammal can be afflicted with a condition such as, *e.g.*, ischemic neuronal damage, traumatic brain injury, peripheral neuropathy, neuropathic pain, Alzheimer's disease, Huntington's disease, Parkinson's disease, amyotrophic lateral sclerosis, and memory impairment. In some embodiments, the mammal is afflicted with a neuronal disorder of the  
5 peripheral nervous system, the medulla, or the spinal cord.

The invention also provides a method of treating a neurodegenerative disease or disorder in a mammal by administering to the mammal a nucleic acid encoding a truncated neublastin polypeptide.

Also provided by the invention is a method of treating a neurodegenerative disease or  
10 disorder in an animal by administering to the animal the truncated neublastin polypeptide. In another aspect, the invention features a method of treating a peripheral neuropathy in a mammal, comprising administering a therapeutically effective amount of a truncated neublastin polypeptide to the mammal. The peripheral neuropathy can be, *e.g.*, one or more of trauma-induced neuropathies, viral-induced neuropathies, chemotherapy-induced neuropathies, toxin-  
15 induced neuropathies, drug-induced neuropathies, vitamin-deficiency-induced neuropathies; idiopathic neuropathies; and diabetic neuropathies. In some embodiments, the truncated neublastin polypeptide is delivered directly into the central nervous system. In other embodiments, the truncated neublastin polypeptide is preferably delivered systemically by subcutaneous injection, intramuscular, intravenous administration, or intravenous infusion.

20 Also within the invention is a method of treating neuropathic pain in a mammal, comprising administering a therapeutically effective amount of a truncated neublastin polypeptide to the mammal. In some embodiments, neuropathic pain associated with toxin-induced nerve damage, pathogen-induced nerve damage, inflammation-induced nerve damage, or neurodegeneration.

25 In a further aspect, the invention features a method of treating a peripheral neuropathy in a mammal by administering a therapeutically effective amount of a nucleic acid encoding truncated neublastin polypeptide to the mammal. The peripheral neuropathy is preferably one or more of trauma-induced neuropathies, chemotherapy-induced neuropathies, toxin-induced neuropathies, viral-induced neuropathies, drug-induced neuropathies, vitamin-deficiency-  
30 induced neuropathies; idiopathic neuropathies; and diabetic neuropathies. Preferably, the nucleic acid encoding the truncated neublastin polypeptide is delivered directly into the central nervous system. Preferably, the nucleic acid encoding the truncated neublastin polypeptide is

delivered systemically by subcutaneous injection, intravenous administration, or intravenous infusion.

Also provided by the invention is a kit that includes, in one or more containers, a substance selected from the group consisting of a truncated neublastin polypeptide and a  
 5 nucleic acid encoding a truncated neublastin polypeptide.

A "neublastin polypeptide," as used herein, is a polypeptide that possesses neurotrophic activity (*e.g.*, as described in Examples 6, 7, 8, and 9) and includes those polypeptides which have an amino acid sequence that has at least 70% homology to the human "neublastin" polypeptides set forth in AA<sub>95</sub>-AA<sub>105</sub> of SEQ ID NO:2, AA<sub>1</sub>-AA<sub>105</sub> of SEQ ID NO:2, AA<sub>97</sub>-AA<sub>140</sub> of SEQ  
 10 ID NO:4, AA<sub>41</sub>-AA<sub>140</sub> of SEQ ID NO:4 ("pro"), AA<sub>1</sub>-AA<sub>140</sub> of SEQ ID NO:4, AA<sub>80</sub>-AA<sub>140</sub> of SEQ ID NO:9 ("wild type" preproNBN), AA<sub>41</sub>-AA<sub>140</sub> of SEQ ID NO:9 (proNBN), AA<sub>1</sub>-AA<sub>140</sub> of SEQ ID NO:5 (mature 140AA), AA<sub>1</sub>-AA<sub>116</sub> of SEQ ID NO:6 (mature 116AA), AA<sub>1</sub>-AA<sub>113</sub> of SEQ ID NO:7 (mature 113AA), AA<sub>1</sub>-AA<sub>140</sub> of SEQ ID NO:10 (mature 140AA), AA<sub>1</sub>-AA<sub>116</sub> of SEQ ID NO:11 (mature 116AA), AA<sub>1</sub>-AA<sub>113</sub> of SEQ ID NO:12 (mature 113AA), and  
 15 the truncated polypeptides of SEQ ID NOs:35-48 variants and derivatives thereof. In addition, this invention contemplates those polypeptides that have an amino acid sequence that has at least 70% homology to the murine "neublastin" polypeptides set forth in AA<sub>1</sub>-AA<sub>224</sub> of SEQ ID NO:16 or to the rat neublastin polypeptides set forth in AA<sub>1</sub>-AA<sub>224</sub> of SEQ ID NO:34.

Preferably, the C-terminal sequence of the above identified neublastin polypeptides has  
 20 an amino acid sequence as set forth in AA<sub>72</sub>-AA<sub>105</sub> of SEQ ID NO:2 (*i.e.*, AA<sub>107</sub>-AA<sub>140</sub> of SEQ ID NO:9), more preferably AA<sub>41</sub>-AA<sub>105</sub> of SEQ ID NO:2 (*i.e.*, AA<sub>76</sub>-AA<sub>140</sub> of SEQ ID NO:9), or the amino acid sequence set forth in AA<sub>191</sub>-AA<sub>224</sub> of SEQ ID NO:16 or 34.

Also, it is preferable that the neublastin polypeptide retain the seven conserved Cys residues that are characteristic of the GDNF family and of the TGF-beta super family. The  
 25 seven conserved cysteine residues are located at positions 16, 43, 47, 80, 81, 109, and 111 of the neublastin polypeptide of SEQ ID NO:12 (mature 113AA). These correspond, *e.g.*, to positions 43, 70, 74, 107, 108, 136 and 138 of SEQ ID NO:9, or, *e.g.*, at positions 127, 154, 158, 191, 192, 220 and 222 of SEQ ID NOs:16 or 34.

Preferably, the neublastin polypeptide has an amino acid sequence with greater than  
 30 85% homology, more preferably with greater than 90% homology, more preferably with greater than 95% homology, most preferably with greater than 99% homology, to the foregoing sequences (*i.e.*, AA<sub>95</sub>-AA<sub>105</sub> of SEQ ID NO:2, AA<sub>1</sub>-AA<sub>105</sub> of SEQ ID NO:2, AA<sub>97</sub>-AA<sub>140</sub> of

SEQ ID NO:4, AA<sub>1</sub>-AA<sub>140</sub> of SEQ ID NO:4, AA<sub>41</sub>-AA<sub>140</sub> of SEQ ID NO:4, AA<sub>80</sub>-AA<sub>140</sub> of  
 SEQ ID NO:9 ("wild type" prepro), AA<sub>41</sub>-AA<sub>140</sub> of SEQ ID NO:9 (pro), AA<sub>1</sub>-AA<sub>140</sub> of SEQ  
 ID NO:5 (mature 140AA), AA<sub>1</sub>-AA<sub>116</sub> of SEQ ID NO:6 (mature 116AA), AA<sub>1</sub>-AA<sub>113</sub> of SEQ  
 ID NO:7 (mature 113AA), AA<sub>1</sub>-AA<sub>140</sub> of SEQ ID NO:10 (mature 140AA), AA<sub>1</sub>-AA<sub>116</sub> of SEQ  
 5 ID NO:11 (mature 116AA), AA<sub>1</sub>-AA<sub>113</sub> of SEQ ID NO:12 (mature 113AA), AA<sub>1</sub>-AA<sub>224</sub> of  
 SEQ ID NO:16 or 34, and SEQ ID NOS:35, 36, 37, 38, 39, 40, 41, 42, 43, 44, 45, 46, 47 and  
 48 (truncated NBN112 through NBN99, respectively).

A "neublastin nucleic acid," as used herein, is a polynucleotide which codes for a  
 neublastin polypeptide. Accordingly, an isolated neublastin nucleic acid is a polynucleotide  
 10 molecule having an open reading frame of nucleotide codons that, were it to be exposed to the  
 appropriate components required for translation, would encode, or code for, a neublastin  
 polypeptide. Neublastin nucleic acids of the invention may be RNA or DNA, *e.g.*, genomic  
 DNA, or DNA which is complementary to and/or transcribed from, a neublastin mRNA  
 ("cDNA"). Thus, a neublastin nucleic acid of the invention further includes polynucleotide  
 15 molecules which hybridize with specificity, under high stringency hybridization conditions, to a  
 polynucleotide that codes for a neublastin polypeptide. This invention also relates to nucleic  
 acid primers that are useful in identifying, isolating and amplifying polynucleotides that encode  
 neublastin polypeptides, or fragments thereof. In certain embodiments of the invention,  
 certain of these primers are neublastin-specific probes useful for hybridization to a neublastin  
 20 nucleic acid, but not to nucleic acids coding for the other members of the GDNF family. By  
 "specific", "specificity", or "specifically", is meant an ability to hybridize with neublastin  
 nucleic acid and inability to hybridize with non-neublastin nucleic acids, including an inability  
 to hybridize to nucleic acids that code uniquely for the other GDNF ligands (*e.g.*, GDNF,  
 persephin, and neurturin).

25 In another embodiment, a neublastin nucleic acid of the invention is one that is  
 identified as being complementary to a polynucleotide that codes for a neublastin polypeptide,  
 either by having a complementary nucleic acid sequence or demonstrating that it hybridizes  
 with specificity at high stringency hybridization conditions to a polynucleotide that codes for  
 neublastin. Particular neublastin nucleic acids include, without limitation, the nucleic acid  
 30 sequences shown herein and designated SEQ ID NO:1, SEQ ID NO:3, SEQ ID NO:8, SEQ ID  
 NO:13, SEQ ID NO:14, SEQ ID NO:15, SEQ ID NO:29 and SEQ ID NO:30 as well as primers  
 SEQ ID NOS: 17-28, 31 and 32. A neublastin nucleic acid of the invention further includes a



unique subregion, or fragment, of a neublastin nucleic acid, including without limitation the nucleic acid fragments shown in FIG. 8.

The neublastin nucleic acids of the invention may be used to express a neublastin polypeptide, *e.g.*, by expressing a neublastin polypeptide *in vivo*, or by administering a  
5 neublastin nucleic acid to an animal for *in vivo* expression. Neublastin nucleic acids may be included within a nucleic acid vector, *e.g.*, an expression vector or a cloning vector. A neublastin nucleic acid may, but need not of necessity, be maintained, reproduced, transferred, or expressed as part of a nucleic acid vector. A recombinant expression vector containing a neublastin polynucleotide sequence can be introduced into and/or maintained within a cell.  
10 Cells hosting a neublastin vector may be prokaryotic. Alternatively, a neublastin nucleic acid can be introduced into a eukaryotic cell, *e.g.*, a eukaryotic cell that contains the appropriate apparatus for post-translational processing of a polypeptide into a mature protein, and/or the appropriate apparatus for secreting a polypeptide into the extracellular environment of the cell.

The invention further features a neublastin neurotrophic factor, "neublastin." Neublastin  
15 may be in the form of a polypeptide, or may be a multimer of two or more neublastin polypeptides, *e.g.*, a neublastin dimer. Neublastin polypeptides are associated as multimers by intermolecular structural associations known to those skilled in the art, including without limitation cysteine-cysteine interaction, disulfide bonds, and noncovalent interactions. Particular neublastin polypeptides include, without limitation, an amino acid sequence disclosed herein  
20 and designated SEQ ID NO:2; SEQ ID NO:4; SEQ ID NO:5; SEQ ID NO:6; SEQ ID NO:7; SEQ ID NO:9; SEQ ID NO:10; SEQ ID NO:11, SEQ ID NO:12, SEQ ID NO:16, SEQ ID NO:34 and SEQ ID NOS:35-48. Preferably, the neublastin polypeptides of the present invention, when dimerized, binds to RET. More preferably, the present neublastin polypeptides, when dimerized, induce dimerization of RET. RET dimerization on the surface  
25 of a cell leads to autophosphorylation of the RET dimer and ultimately to the activation of the RET mediated intracellular signaling cascade.

A neublastin polypeptide of the invention is useful for treating a defect in a neuron, including without limitation lesioned neurons and traumatized neurons. Peripheral nerves that experience trauma include, but are not limited to, nerves of the medulla or of the spinal cord.  
30 Neublastin polypeptides are useful in the treatment of neurodegenerative disease, *e.g.*, cerebral ischemic neuronal damage; neuropathy, *e.g.*, peripheral neuropathy, Alzheimer's disease, Huntington's disease, Parkinson's disease, amyotrophic lateral sclerosis (ALS). Neublastin

polypeptides are further contemplated for use in the treatment of impaired memory, *e.g.*, memory impairment associated with dementia.

Additional examples of conditions or diseases are disorders of the peripheral nervous system, the medulla, or the spinal cord, as well as trauma-induced neuropathies, chemotherapy-  
5 induced neuropathies, toxin-induced neuropathies, drug-induced neuropathies, vitamin-deficiency-induced neuropathies; idiopathic neuropathies; and diabetic neuropathies, neuropathic pain associated with toxin-induced nerve damage, pathogen-induced nerve damage, inflammation-induced nerve damage, or neurodegeneration. Additional examples of peripheral neuropathies include trauma-induced neuropathies, chemotherapy-induced  
10 neuropathies, toxin-induced neuropathies, drug-induced neuropathies, vitamin-deficiency-induced neuropathies; idiopathic neuropathies; and diabetic neuropathies.

Unless otherwise defined, all technical and scientific terms used herein have the same meaning as commonly understood by one of ordinary skill in the art to which this invention belongs. Although methods and materials similar or equivalent to those described herein can  
15 be used in the practice or testing of the invention, suitable methods and materials are described below. All publications, patent applications, patents, and other references mentioned herein are incorporated by reference in their entirety. In the case of conflict, the present specification, including definitions, will control. In addition, the materials, methods, and examples are illustrative only and not intended to be limiting.

20 Other features and advantages of the invention will be apparent from the following detailed description and claims.

### BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1 is a photographic image of two northern blots probed with <sup>32</sup>P-labelled neublastin  
25 cDNA, comparing relative levels of expression of the neublastin gene in various human adult tissue types (panel A) and in various regions of the adult human brain (panel B).

FIG. 2 is a photographic image of a northern blot probed with <sup>32</sup>P-labelled neublastin cDNA, comparing the amount of neublastin cDNA expressed in a non-transfected cell-line, HiB5, with the amount of neublastin cDNA expressed in a cell-line transfected with neublastin  
30 cDNA, and with a cell-line transfected with GDNF- cDNA.

FIG. 3 is a photographic image of two western blots which compare the degrees to which neublastin protein is expressed in non-transfected HiB5 cells (lane 1) relative to an HiB5

cell-line stably-transfected with neublastin cDNA (lane 2) was probed with either the neublastin-specific antibody Ab-2 (left blot; Panel A) or the neublastin-specific antibody Ab-1 (right blot; Panel B).

FIG. 4 is a graphical illustration of the effect of neublastin on the survival of cultured rat embryonic, dopaminergic, ventral mesencephalic neurons and ChAT activity in cholinergic cranial nerve motor neurons in serum-free medium. In particular, FIG. 4A is an illustration of the dose-response curve for recombinant GDNF on ChAT activity (dpm/hour). FIG. 4B is an illustration of ChAT activity (dpm/hour) using diluted conditioned medium from either neublastin producing or GDNF-producing cells. FIG. 4C is an illustration of the number of tyrosine hydroxylase immunoreactive cells per well.

FIG. 5 is an illustration of the effect of neublastin secreted from HiB5pUbi1zNBN22 cells on the function and survival of slice cultures of pig embryonic dopaminergic ventral mesencephalic neurons co-cultured with either HiB5pUbi1zNBN22 cells (neublastin) or HiB5 cells (control). FIG. 5A and FIG. 5B illustrate dopamine released to the medium at DIV12 (Dopamine (pmol/ml) - day 12) and DIV21 (Dopamine (pmol/ml) - day 21), respectively. FIG. 5C is an illustration of the number of tyrosine hydroxylase immunoreactive cells per culture (TH-ir cells per culture) at DIV21.

FIG. 6 is an illustration of the *in vivo* effect of lentiviral-produced neublastin on nigral dopamine neurons.

FIG. 7 is a schematic diagram of the genomic structure of the neublastin gene, including the nucleic acid primers which can be used to identify the full length neublastin gene, and their spatial orientation in relation to the genomic Neublastin-encoding sequence (*i.e.*, gene).

FIG. 8 is an illustration of neublastin specific primers used to identify the cDNA clone encoding the human neublastin polypeptide that hybridize to nucleic acids that encode neublastin polypeptides, but do not hybridize to nucleic acids encoding the other known GDNF family members (*i.e.*, GDNF, Persephin and neurturin).

FIG. 9 illustrates the neurotrophic activity on cultures of dissociated rat dorsal root ganglion cells from different development stages of a polypeptide disclosed in the present invention in comparison to those obtained with known neurotrophic factors (0 control experiment (in absence of factors); 1 in the presence of GDNF; 2 in the presence of Neurturin; 3 in the presence of Neublastin of the invention; E12 embryonic day 12; E16 embryonic day 16; P0 the day of birth; P7 day 7 after birth; and P15 day 15 after birth).

FIG. 10 illustrates neublastin production from CHO cell lines.

FIG. 11 illustrates a comparison of neublastin and GDNF binding to GFR $\alpha$ -1 and GFR $\alpha$ -3 receptors.

FIG. 12 is a photographic image of a western blot which illustrates R30 anti-peptide  
5 antibody and R31 anti-peptide antibody binding to neublastin.

FIG. 13 is a picture of a gel showing extraction of neublastin by affinity binding on RETL3-Ig.

FIG. 14 is a plasmid map of pET19b-Neublastin, along with the sequence of the synthetic gene for Neublastin.

10 FIG. 15 is a plasmid map of pMJB164-HisNeublastin, along with the sequence of the synthetic gene for HisNeublastin.

FIG. 16 illustrates a comparison of a 102 amino acid form of truncated neublastin (NBN) and 113 amino acid form of neublastin in a cellular RET Activation assay.

FIG. 17 illustrates a comparison of various forms of neublastin or neublastin muteins in  
15 a cellular RET Activation assay.

FIG. 18 illustrates a comparison of various mature (113 aa) and truncated (106 aa, 104 aa or 99 aa) forms of neublastin or neublastin R14K muteins in a cellular KIRA ELISA assay.

#### DETAILED DISCLOSURE OF THE INVENTION

Applicants have identified a nucleic acid that encodes a novel neurotrophic factor that is  
20 referred to herein as "neublastin," or "NBN." Neublastin is a member of the glial cell line-derived neurotrophic factor (GDNF) sub-class of the transforming growth factor- $\beta$  (TGF- $\beta$ ) super-family of neurotrophic factors. NBN polypeptides are biologically active as dimers, and have little to no neurotrophic activity as monomers. Therefore, reference herein to any neublastin polypeptide of the invention is understood to designate the homodimeric form of the designated  
25 neublastin, unless specified otherwise.

The cDNA encoding neublastin was originally identified as follows. Using the TBLASTN 1.4.11 algorithm (Atschul et al., *Nucl. Acids Res.*, 25, pp. 3389-3402 (1997)) and human persephin as query (GenBank Acc. No. AF040962), a 290 bp fragment was initially identified in High-Throughput Genomic Sequence (HTGS) of two human bacterial artificial  
30 chromosomes (BAC) with GenBank entries AC005038 and AC005051. AC005038 consists of approximately 190,000 bp of 5 contigs of unordered sequences and AC005051 consists of approximately 132,000 bp of 12 contigs of unordered sequences. The 290 bp fragment

identified in the two BAC clones proved to have regions that were homologous, but not identical, to a coding region of the cDNA of the neurotrophic factor, human persephin.

From this 290 bp sequence two Neublastin-specific PCR primers were synthesised (Top Stand Primer (SEQ ID NO:17) and Bottom Strand Primer (SEQ ID NO:18)). Screening of human fetal brain cDNA library was performed. The initial screening comprised 96-well PCR-based screening with the two PCR primers (SEQ ID NOS. 17 and 18) of a cDNA library "Master Plate" from 500,000 cDNA clones containing approximately 5,000 clones/well. A second PCR-based screen was performed on a human fetal brain cDNA library "Sub-Plate" containing *E. coli* glycerol stock with approximately 5,000 clones/well.

A 102 bp fragment (SEQ ID NO:13) was identified in the PCR-based screenings of both the Master Plate and Sub Plate. A positive cDNA clone (possessing the 102 bp fragment) was selected, plated on two LB/antibiotic-containing plates, and grown overnight. From these plates, a total of 96 bacterial colonies were selected and individually placed in the wells of a new, 96-well PCR plate containing both PCR primers (SEQ ID NOS. 17 and 18) and the requisite PCR amplification reagents. PCR amplification was then performed and the 96 individual PCR reactions were analyzed by 2% agarose gel electrophoresis. The positive colony with the clone containing the 102 bp fragment was then identified. Plasmid DNA was obtained from the positive colony containing the 102 bp fragment and sequenced. Subsequent sequencing analysis revealed the presence of a full-length cDNA of 861 bp (SEQ ID NO:8). The Open Reading Frame (ORF) of 663 bp, also referred to as the coding region (CDS), identified in SEQ ID NO:8, encodes the pre-pro-polypeptide (designated "pre-pro-Neublastin") and is shown in SEQ ID NO:9. Based on SEQ ID NO:9, three variants of Neublastin polypeptides were identified. These variants include:

- (i) the 140 AA polypeptide designated herein as NBN140, which possesses the amino acid sequence designated as SEQ ID NO:10;
- (ii) the 116 AA polypeptide designated herein as NBN116, which possesses the amino acid sequence designated as SEQ ID NO:11; and
- (iii) the 113 AA polypeptide designated herein as NBN113, which possesses the amino acid sequence designated as SEQ ID NO:12.

Other variants of Neublastin include truncated NBN forms. Examples of these include:

- (iv) the 112AA polypeptide sequence designated herein as NBN112, which possesses the 112 carboxy terminal amino acids of a neublastin polypeptide, *e.g.*, amino acids 29-140 of SEQ ID NO:9 (SEQ ID NO:35) or amino acids 113-224 of SEQ ID NOs:16 or 34.
- 5 (v) the 111AA polypeptide sequence designated herein as NBN111, which possesses the 111 carboxy terminal amino acids of a neublastin polypeptide, *e.g.*, amino acids 30-140 of SEQ ID NO:9 (SEQ ID NO:36) or amino acids 114-224 of SEQ ID NOs:16 or 34.
- 10 (vi) the 110AA polypeptide sequence designated herein as NBN110, which possesses the 110 carboxy terminal amino acids of a neublastin polypeptide, *e.g.*, amino acids 31-140 of SEQ ID NO:9 (SEQ ID NO:37) or amino acids 115-224 of SEQ ID NOs:16 or 34.
- 15 (vii) the 109AA polypeptide sequence designated herein as NBN109, which possesses the 109 carboxy terminal amino acids of a neublastin polypeptide, *e.g.*, amino acids 32-140 of SEQ ID NO:9 (SEQ ID NO:38) or amino acids 116-224 of SEQ ID NOs:16 or 34.
- 20 (viii) the 108AA polypeptide sequence designated herein as NBN108, which possesses the 108 carboxy terminal amino acids of a neublastin polypeptide, *e.g.*, amino acids 33-140 of SEQ ID NO:9 (SEQ ID NO:39) or amino acids 117-224 of SEQ ID NOs:16 or 34.
- (ix) the 107AA polypeptide sequence designated herein as NBN107, which possesses the 107 carboxy terminal amino acids of a neublastin polypeptide, *e.g.*, amino acids 34-140 of SEQ ID NO:9 (SEQ ID NO:40) or amino acids 118-224 of SEQ ID NOs:16 or 34.
- 25 (x) the 106AA polypeptide sequence designated herein alternatively as designated herein as NBN106 or N-7, which possesses the 106 carboxy terminal amino acids of a neublastin polypeptide, *e.g.*, amino acids 35-140 of SEQ ID NO:9 (SEQ ID NO:41) or amino acids 119-224 of SEQ ID NOs:16 or 34.

- (xi) the 105AA polypeptide sequence designated herein as NBN105, which possesses the 105 carboxy terminal amino acids of a neublastin polypeptide, *e.g.*, amino acids 36-140 of SEQ ID NO:9 (SEQ ID NO:42) or amino acids 120-224 of SEQ ID NOs:16 or 34.
- 5 (xii) the 104AA polypeptide sequence designated herein alternatively as designated herein as NBN104 or N-9, which possesses the 104 carboxy terminal amino acids of a neublastin polypeptide, *e.g.*, amino acids 37-140 of SEQ ID NO:9 (SEQ ID NO:43) or amino acids 121-224 of SEQ ID NOs:16 or 34.
- 10 (xiii) the 103AA polypeptide sequence designated herein as NBN103, which possesses the 103 carboxy terminal amino acids of a neublastin polypeptide, *e.g.*, amino acids 38-140 of SEQ ID NO:9 (SEQ ID NO:44) or amino acids 122-224 of SEQ ID NOs:16 or 34.
- 15 (xiv) the 102AA polypeptide sequence designated herein as NBN102, which possesses the 102 carboxy terminal amino acids of a neublastin polypeptide, *e.g.*, amino acids 39-140 of SEQ ID NO:9 (SEQ ID NO:45) or amino acids 123-224 of SEQ ID NOs:16 or 34.
- 20 (xv) the 101AA polypeptide sequence designated herein as NBN101, which possesses the 101 carboxy terminal amino acids of a neublastin polypeptide, *e.g.*, amino acids 40-140 of SEQ ID NO:9 (SEQ ID NO:46) or amino acids 124-224 of SEQ ID NOs:16 or 34.
- (xvi) the 100AA polypeptide sequence designated herein as NBN100, which possesses the 100 carboxy terminal amino acids of a neublastin polypeptide, *e.g.*, amino acids 41-140 of SEQ ID NO:9 (SEQ ID NO:47) or amino acids 125-224 of SEQ ID NOs:16 or 34.
- 25 (xvii) the 99AA polypeptide sequence designated herein alternatively as NBN99 or N-14, which possesses the 99 carboxy terminal amino acids of a neublastin polypeptide, *e.g.*, amino acids 42-140 of SEQ ID NO:9 (SEQ ID NO:48) or amino acids 126-224 of SEQ ID NOs:16 or 34.

The neublastin polypeptides described herein may be provided in any bioactive form,  
30 including the form of pre-pro-proteins, pro-proteins, mature proteins, glycosylated proteins,

non-glycosylated proteins, phosphorylated proteins, non-phosphorylated proteins, truncated forms, or any other post-translationally modified protein. It is assumed that a bioactive neublastin is in the dimerized form for each NBN variant, where dimer formation is required for activity. Little to no neurotrophic activity is observed in a monomeric NBN polypeptide. A  
5 bioactive neublastin polypeptide includes a dimerized polypeptide that, in the presence of a cofactor (such as GFR $\alpha$ 3 or RET), binds to GFR $\alpha$ 3 or to a complex of GFR $\alpha$ 3 and RET, induces dimerization of RET, and autophosphorylation of RET. It is understood that the truncated forms of Neublastin disclosed herein (e.g., the 112AA, 111AA, 110AA, 109AA, 108AA, 107AA, 106AA, 105AA, 104AA, 103AA, 102AA, 101AA, 100AA or 99AA forms,  
10 shown in SEQ ID NOS:35-48, respectively), as well as all other NBN polypeptides described above, are dimers, and that each dimer has neurotrophic activity.

The entire cDNA sequence containing 782 bp 5' untranslated DNA, 663 bp encoding DNA, and 447 3' untranslated (totalling 1992 bp) has been submitted to GenBank under the Accession Number AF 120274.

15 The genomic Neublastin-encoding sequence was identified as follows:

With the goal of cloning the genomic neublastin-encoding sequence, an additional set of primers were prepared. In particular, Primer Pair No.1 comprised (sense shown as SEQ ID NO:23 and antisense shown as SEQ ID NO:24) and Primer Pair No. 2 comprised (sense shown as SEQ ID NO:25 and antisense shown as SEQ ID NO:26).

20 Using Primer Pair No. 2, a 887 bp DNA fragment was amplified by PCR from a preparation of human genomic DNA, and cloned into the pCRII vector (Invitrogen) and transformed into *E. coli*. The resulting plasmid was sequenced and a 861 bp putative cDNA sequence (encoding a protein named neublastin herein) was predicted (as set forth in SEQ ID NO:3). Similarly, using Primer Pair No. 1, an 870 bp DNA fragment was obtained by PCR of  
25 human genomic DNA. An additional 42 bp region at the 3'-terminus of the Open Reading Frame (ORF) was found in this fragment, in comparison to the 887 bp sequence. The genomic structure of the neublastin gene was predicted by comparing it to the sequences of nucleic acids of other neurotrophic factors, by mapping exon-intron boundaries. This analysis demonstrated that the neublastin gene has at least two exons separated by a 70 bp intron.

30 This sequence was also used to screen GenBank for neublastin EST sequences. Three were identified with GenBank entries AA844072, AA931637 and AA533512, indicating that neublastin nucleic acids are transcribed into mRNA.



Comparison of the entire cDNA sequence obtained (AF 120274) and the genomic sequence present in GenBank entries AC005038 and AC005051 revealed that the neublastin gene consists of at least five exons (including three coding) separated by four introns (see, *e.g.*, FIG. 7). Together, the exons have a predicted amino acid sequence of a full-length Neublastin polypeptide. It should also be noted that the 887 bp fragment was found to contain the complete coding region of pro-neublastin. The predicted cDNA (SEQ ID NO:3) contains an Open Reading Frame (ORF) encoding pro-neublastin (181 amino acid residues) which showed homology to the three known human proteins - Persephin, Neurturin, and GDNF. See Table 3 in Examples.

#### 10        **Neublastin Nucleic Acids of the Invention**

In another aspect, the invention provides polynucleotides capable of expressing the polypeptides of the invention. The polynucleotides of the invention include DNA, cDNA and RNA sequences, as well as anti-sense sequences, and include naturally occurring, synthetic, and intentionally manipulated polynucleotides. The polynucleotides of the invention also include sequences that are degenerate as a result of the genetic code, but which code on expression for a neublastin polypeptide.

As defined herein, the term "polynucleotide" refers to a polymeric form of nucleotides of at least 10 bases in length, preferably at least 15 bases in length. By "isolated polynucleotide" is meant a polynucleotide that is not immediately contiguous with both of the coding sequences with which it is immediately contiguous (one on the 5' end and one on the 3' end) in the naturally occurring genome of the organism from which it is derived. The term therefore includes recombinant DNA which is incorporated into an expression vector, into an autonomously replicating plasmid or virus, or into the genomic DNA of a prokaryote or eukaryote, or which exists as a separate molecule, *e.g.* a cDNA, independent from other sequences.

The polynucleotides of the invention also include allelic variants and "mutated polynucleotides" having a nucleotide sequence that differs from the nucleotide sequences presented herein at one or more nucleotide positions.

In a preferred embodiment, the polynucleotide of the invention has a nucleic acid (DNA) sequence capable of hybridizing with the polynucleotide sequence presented as SEQ ID NO:1, the polynucleotide sequence presented as SEQ ID NO:3, the polynucleotide sequence presented as SEQ ID NO:8, or the polynucleotide sequence presented as SEQ ID NO:15, its

complementary strand, or a sub-sequence hereof under at least medium, medium/high, or high stringency conditions, as described in more detail below.

In another preferred embodiment, the isolated polynucleotide of the invention has a nucleic acid (DNA) sequence that is at least 70%, preferably at least 80%, more preferred at  
5 least 90%, more preferred at least 95%, most preferred at least 99% homologous to the polynucleotide sequence presented as SEQ ID NO:1, the polynucleotide sequence presented as SEQ ID NO:3, the polynucleotide sequence presented as SEQ ID NO:8, or the polynucleotide sequence presented as SEQ ID NO:15.

In its most preferred embodiment, the polynucleotide has the DNA sequence presented  
10 as SEQ ID NO:1, the DNA sequence presented as SEQ ID NO:3, the DNA sequence presented as SEQ ID NO:8, or the polynucleotide sequence presented as SEQ ID NO:15.

This invention also provides novel primers and DNA sequences for identifying, isolating and amplifying neublastin polynucleotides which code on expression for neublastin polypeptides or fragments thereof. Such primers include the polynucleotides set forth in SEQ  
15 ID NOS:17-28, and 31-32. In addition, this invention provides neublastin DNA sequences generated from those primers, including those set forth in SEQ ID NOS:13 and 14. Further, this invention provides DNA sequences from 3' or 5' untranslated regions ("UTR") in genomic DNA that flank neublastin exons; such sequences are useful in identifying, isolating and amplifying neublastin polynucleotides which code on expression for neublastin polypeptides or  
20 fragments thereof.

3' UTR sequences of this invention include the sequences set forth in:

nucleotides 721 - 865 of SEQ ID NO:1,  
nucleotides 718 - 861 of SEQ ID NO:3,  
nucleotides 718 - 861 of SEQ ID NO:8,  
25 nucleotides 1647 - 2136 of SEQ ID NO:15, and  
contiguous sequences of between 10 - 25 nucleotides derived from (*i.e.*, falling within) the foregoing sequences (which are useful, *e.g.*, as primers).

5' UTR sequences of this invention include the sequences set forth in:

nucleotides 1-10 of SEQ ID NO:1,  
30 nucleotides 1 - 57 of SEQ ID NO:8,  
nucleotides 1-974 of SEQ ID NO:15, and

contiguous sequences of between 10 - 25 nucleotides derived from (*i.e.*, falling within) the foregoing sequences (which are useful, *e.g.*, as primers).

The polynucleotides of the invention may preferably be obtained by cloning procedures, *e.g.* as described in "Current Protocols in Molecular Biology" (John Wiley & Sons, Inc.). In a preferred embodiment, the polynucleotide is cloned from, or produced on the basis of human genomic DNA or a cDNA library of the human brain.

#### **Homology of DNA sequences**

The DNA sequence homology referred to above may be determined as the degree of identity between two sequences indicating a derivation of the first sequence from the second.

The homology may suitably be determined by means of computer programs known in the art, such as GAP provided in the GCG program package (*Needleman, S.B. and Wunsch C.D., Journal of Molecular Biology* 1970 **48**, 443-453). Using GAP with the following settings for DNA sequence comparison: GAP creation penalty of 5.0 and GAP extension penalty of 0.3, the coding region of the analogous DNA sequences referred to above exhibits a degree of identity preferably of at least 70%, more preferably at least 80%, more preferably at least 90%, more preferably at least 95%, most preferably at least 99%, with the CDS (encoding) part of the DNA sequence shown in SEQ ID NO:1, or the CDS (encoding) part of the DNA sequence shown in SEQ ID NO:3, or the CDS (encoding) part of the DNA sequence shown in SEQ ID NO:8, the CDS (encoding) part of the DNA sequence shown in SEQ ID NO:15.

The term "sequence identity" refers to the degree to which two polynucleotide sequences are identical on a nucleotide-by-nucleotide basis over a particular region of comparison. The term "percentage of sequence identity" is calculated by comparing two optimally aligned sequences over that region of comparison, determining the number of positions at which the identical nucleic acid base (*e.g.*, A, T, C, G, U, or I) occurs in both sequences to yield the number of matched positions, dividing the number of matched positions by the total number of positions in the region of comparison (*i.e.*, the window size), and multiplying the result by 100 to yield the percentage of sequence identity. The term "substantial identity" as used herein denotes a characteristic of a polynucleotide sequence, wherein the polynucleotide comprises a sequence that has at least 80 percent sequence identity, preferably at least 85 percent identity and often 90 to 95 percent sequence identity, more usually at least 99 percent sequence identity as compared to a reference sequence over a

comparison region. Methods of determining sequence identity are well known in the art. See, e.g., Needleman and Wunsch J. Mol. Biol. 1970 **48**, 443-453.

#### **Hybridization Protocol**

5 The polynucleotides of the invention are such which have a nucleic acid sequence capable of hybridizing with the polynucleotide sequence presented as SEQ ID NO:1, the polynucleotide sequence presented as SEQ ID NO:3, or the polynucleotide sequence presented as SEQ ID NO:8, or the polynucleotide sequence presented as SEQ ID NO:15, or their complementary strand, or a sub-sequence hereof under at least medium, medium/high, or high stringency conditions, as described in more detail below.

10 Suitable experimental conditions for determining hybridization between a nucleotide probe and a homologous DNA or RNA sequence, involves pre-soaking of the filter containing the DNA fragments or RNA to hybridize in 5 X SSC (Sodium chloride/Sodium citrate; cf. *Sambrook et al.*; Molecular Cloning: A Laboratory Manual, Cold Spring Harbor Lab., Cold Spring Harbor, NY 1989) for 10 minutes, and pre-hybridization of the filter in a solution of 5 X  
15 SSC, 5 X Denhardt's solution (cf. *Sambrook et al.*; *Op cit.*), 0.5 % SDS and 100 µg/ml of denatured sonicated salmon sperm DNA (cf. *Sambrook et al.*; *Op cit.*), followed by hybridization in the same solution containing a concentration of 10 ng/ml of a random-primed (*Feinberg A P & Vogelstein B*; Anal. Biochem. 1983 **132** 6-13), <sup>32</sup>P-dCTP-labeled (specific activity > 1 x 10<sup>9</sup> cpm/µg) probe for 12 hours at approximately 45°C. The filter is then washed  
20 twice for 30 minutes in 0.1 x SSC, 0.5 % SDS at a temperature of at least at least 60°C (medium stringency conditions), preferably of at least 65°C (medium/high stringency conditions), more preferred of at least 70°C (high stringency conditions), and even more preferred of at least 75°C (very high stringency conditions). Molecules to which the oligonucleotide probe hybridizes under these conditions may be detected using a X-ray film.

#### **Cloned Polynucleotides**

25 The isolated polynucleotide of the invention may in particular be a cloned polynucleotide. As defined herein, the term "cloned polynucleotide" refers to a polynucleotide or DNA sequence cloned in accordance with standard cloning procedures currently used in genetic engineering to relocate a segment of DNA, which may in particular be cDNA, i.e.,  
30 enzymatically derived from RNA, from its natural location to a different site where it will be reproduced. The cloned polynucleotide of the invention can encode any one of the NBN

polypeptides disclosed herein, including but not limited to the polypeptides shown in SEQ ID NOS: 2, 4-7, 9-12, 16, 34, and 35-48.

- Cloning may be accomplished by any suitable route and may involve techniques such as reverse transcriptase technology, PCR technology, and the like, as well as excision and
- 5 isolation of the desired DNA segment.

The cloned polynucleotide of the invention may alternatively be termed "DNA construct" or "isolated DNA sequence", and may in particular be a complementary DNA (cDNA).

#### **Biological Sources**

- 10 The isolated polynucleotide of the invention may be obtained from any suitable source.

In a preferred embodiment, which the polynucleotide of the invention is cloned from, or produced on the basis of a cDNA library, *e.g.* of a cDNA library of the fetal or adult brain, in particular of the forebrain, the hindbrain, the cortex, the striatum, the amygdala, the cerebellum, the caudate nucleus, the corpus callosum, the hippocampus, the thalamic nucleus,

15 the subthalamic nucleus, the olfactory nucleus, the putamen, the substantia nigra, the dorsal root ganglia, the trigeminal ganglion, the superior mesenteric artery, or the thalamus; of the spinal cord; of the heart; the placenta; of the lung; of the liver; of the skeletal muscle; of the kidney; of the liver; of the pancreas; of the intestines; of the eye; of the retina; of the tooth pulp; of the hair follicle; of the prostate; of the pituitary; or of the trachea.

- 20 Commercial cDNA libraries from a variety of tissues, both human and non-human, are available from *e.g.* Stratagene and Clontech. The isolated polynucleotide of the invention may be obtained by standard methods, *e.g.* those described in the working examples.

#### **Neublastin Polypeptides of the Invention**

- As noted above, a "neublastin polypeptide," as used herein, is a polypeptide which
- 25 possesses neurotrophic activity (*e.g.*, as described in Examples 6, 7, 8, and 9) and includes those polypeptides which have an amino acid sequence that has at least 70% homology to the "neublastin" polypeptides set forth in AA<sub>95</sub>-AA<sub>105</sub> of SEQ ID NO:2, AA<sub>1</sub>-AA<sub>105</sub> of SEQ ID NO:2, AA<sub>97</sub>-AA<sub>140</sub> of SEQ ID NO:4, AA<sub>41</sub>-AA<sub>140</sub> of SEQ ID NO:4, AA<sub>1</sub>-AA<sub>140</sub> of SEQ ID NO:4, AA<sub>80</sub>-AA<sub>140</sub> of SEQ ID NO:9 ("wild type" prepro), AA<sub>41</sub>-AA<sub>140</sub> of SEQ ID NO:9
- 30 (pro), AA<sub>1</sub>-AA<sub>140</sub> of SEQ ID NO:5 (mature 140AA), AA<sub>1</sub>-AA<sub>116</sub> of SEQ ID NO:6 (mature 116AA), AA<sub>1</sub>-AA<sub>113</sub> of SEQ ID NO:7 (mature 113AA), AA<sub>1</sub>-AA<sub>140</sub> of SEQ ID NO:10 (mature 140AA, ), AA<sub>1</sub>-AA<sub>116</sub> of SEQ ID NO:11 (mature 116AA), AA<sub>1</sub>-AA<sub>113</sub> of SEQ ID NO:12

(mature 113AA), AA<sub>1</sub>-AA<sub>224</sub> of SEQ ID NO:16 (murine prepro), AA<sub>1</sub>-AA<sub>224</sub> of SEQ ID NO:34 (rat prepro), truncated form NBN112 through NBN99 (SEQ ID NOs:35-48, respectively), and variants and derivatives of each of the foregoing.

Preferably, the C-terminal sequence of the above identified neublastin polypeptides has an amino acid sequence as set forth in AA<sub>72</sub>-AA<sub>105</sub> of SEQ ID NO:2 (*i.e.*, AA<sub>107</sub>-AA<sub>140</sub> of SEQ ID NO:9), more preferably AA<sub>41</sub>-AA<sub>105</sub> of SEQ ID NO:2 (*i.e.*, AA<sub>76</sub>-AA<sub>140</sub> of SEQ ID NO:9).

Also, it is preferable that the neublastin polypeptide retain the seven conserved Cys residues that are characteristic of the GDNF family and of the TGF-beta super family.

Preferably the neublastin polypeptide has an amino acid sequence greater than 85% homology, more preferably greater than 90% homology, more preferably greater than 95% homology, most preferably greater than 99% homology, to the foregoing sequences (*i.e.*, AA<sub>95</sub>-AA<sub>105</sub> of SEQ ID NO:2, AA<sub>1</sub>-AA<sub>105</sub> of SEQ ID NO:2, AA<sub>97</sub>-AA<sub>140</sub> of SEQ ID NO:4, AA<sub>41</sub>-AA<sub>140</sub> of SEQ ID NO:4, AA<sub>1</sub>-AA<sub>140</sub> of SEQ ID NO:4, AA<sub>80</sub>-AA<sub>140</sub> of SEQ ID NO:9 ("wild type" prepro), AA<sub>41</sub>-AA<sub>140</sub> of SEQ ID NO:9 (pro), AA<sub>1</sub>-AA<sub>140</sub> of SEQ ID NO:5 (mature 140AA), AA<sub>1</sub>-AA<sub>116</sub> of SEQ ID NO:6 (mature 116AA), AA<sub>1</sub>-AA<sub>113</sub> of SEQ ID NO:7 (mature 113AA), AA<sub>1</sub>-AA<sub>140</sub> of SEQ ID NO:10 (mature 140AA), AA<sub>1</sub>-AA<sub>116</sub> of SEQ ID NO:11 (mature 116AA), AA<sub>1</sub>-AA<sub>113</sub> of SEQ ID NO:12 (mature 113AA), AA<sub>1</sub>-AA<sub>224</sub> of SEQ ID NO:16 (murine prepro), AA<sub>1</sub>-AA<sub>224</sub> of SEQ ID NO:34 (rat prepro), truncated neublastin polypeptides NBN112 through NBN99 (SEQ ID NOs:35-48, respectively), and preferably any of the foregoing polypeptides with a C-terminal sequence of the above identified neublastin polypeptides has an amino acid sequence as set forth in AA<sub>72</sub>-AA<sub>105</sub> of SEQ ID NO:2 (*i.e.*, AA<sub>107</sub>-AA<sub>140</sub> of SEQ ID NO:9), more preferably AA<sub>41</sub>-AA<sub>105</sub> of SEQ ID NO:2 (*i.e.*, AA<sub>76</sub>-AA<sub>140</sub> of SEQ ID NO:9) or AA<sub>191</sub>-AA<sub>224</sub> of SEQ ID NOs:16 or 34.

In addition, this invention contemplates those polypeptides which have an amino acid sequence that has at least 70% homology to the murine "neublastin" polypeptides set forth in AA<sub>1</sub>-AA<sub>224</sub> of SEQ ID NO:16, or rat neublastin polypeptides set forth in AA<sub>1</sub>-AA<sub>224</sub> of SEQ ID NO:34.

Among the preferred polypeptides of the invention in one embodiment represent the preprosequence (as set forth in SEQ ID NOS:2, 4, 9, 16, and 34, respectively), the pro sequence (as set forth in AA<sub>75</sub>-AA<sub>105</sub> of SEQ ID NO:2, or AA<sub>41</sub>-AA<sub>140</sub> of SEQ ID NOS:4 and 9, respectively) the mature sequence (as set forth in SEQ ID NOS:5, 6, 7, 10, 11, or 12, preferably

SEQ ID NOS:10, 11, 12), and most preferably the truncated sequences (SEQ ID NOS:35-48) of neublastin.

The polypeptides of the invention include variant polypeptides. In the context of this invention, the term "variant polypeptide" means a polypeptide (or protein) having an amino acid sequence that differs from the sequence presented as SEQ ID NO:2, SEQ ID NO:4, SEQ ID NO:5, SEQ ID NO:6, SEQ ID NO:7, SEQ ID NO:9, SEQ ID NO:10, SEQ ID NO:11, SEQ ID NO:12, SEQ ID NO:16, SEQ ID NO:34 or SEQ ID NOS:35-48, at one or more amino acid positions. Such variant polypeptides include the modified polypeptides described above, as well as conservative substitutions, splice variants, isoforms, homologues from other species, and polymorphisms.

As defined herein, the term "conservative substitutions" denotes the replacement of an amino acid residue by another, biologically similar residue. For example, one would expect conservative amino acid substitutions to have little or no effect on the biological activity, particularly if they represent less than 10% of the total number of residues in the polypeptide or protein. Preferably, conservative amino acids substitutions represent changes in less than 5% of the polypeptide or protein, most preferably less than 2% of the polypeptide or protein (*e.g.*, when calculated in accordance with NBN113, most preferred conservative substitutions would represent fewer than 3 amino acid substitutions in the wild type amino acid sequence). In a particularly preferred embodiment, there is a single amino acid substitution in the sequence, wherein the both the substituted and replacement amino acid are non-cyclic.

Other examples of particularly conservative substitutions include the substitution of one hydrophobic residue such as isoleucine, valine, leucine or methionine for another, or the substitution of one polar residue for another, such as the substitution of arginine for lysine, glutamic for aspartic acid, or glutamine for asparagine, and the like.

The term conservative substitution also includes the use of a substituted amino acid residue in place of an un-substituted parent amino acid residue provided that antibodies raised to the substituted polypeptide also immunoreact with the un-substituted polypeptide.

The term "conservative substitution variant" accordingly refers to a neublastin polypeptide which differs from a wild type or reference neublastin polypeptide by the presence of at least one conservative amino acid substitution.

Modifications of a neublastin primary amino acid sequence may result in proteins which have substantially equivalent activity as compared to the unmodified counterpart

polypeptide, and thus may be considered functional analogous of the parent proteins. Such modifications may be deliberate, *e.g.* as by site-directed mutagenesis, or they may occur spontaneous, and include splice variants, isoforms, homologues from other species, and polymorphisms. Such functional analogs are also contemplated according to the invention.

5           Moreover, modifications of the primary amino acid sequence may result in proteins which do not retain the biological activity of the parent protein, including dominant negative forms, etc. A dominant negative protein may interfere with the wild-type protein by binding to, or otherwise sequestering regulating agents, such as upstream or downstream components, that normally interact functionally with the polypeptide. Such dominant negative forms are also  
10           contemplated according to the invention.

A "signal peptide" is a peptide sequence that directs a newly synthesized polypeptide to which the signal peptide is attached to the endoplasmic reticulum (ER) for further post-translational processing and distribution.

15           An "heterologous signal peptide," as used herein in the context of neublastin, means a signal peptide that is not the human neublastin signal peptide, typically the signal peptide of some mammalian protein other than neublastin.

Skilled artisans will recognize that the human neublastin DNA sequence (either cDNA or genomic DNA), or sequences that differ from human neublastin DNA due to either silent codon changes or to codon changes that produce conservative amino acid substitutions, can be  
20           used to genetically modify cultured human cells so that they will overexpress and secrete the enzyme.

Polypeptides of the present invention also include chimeric polypeptides or cleavable fusion polypeptides in which another polypeptide is fused at the N-terminus or the C-terminus of the polypeptide or fragment thereof. A chimeric polypeptide may be produced by fusing a nucleic  
25           acid sequence (or a portion thereof) encoding another polypeptide to a nucleic acid sequence (or a portion thereof) of the present invention.

Techniques for producing chimeric polypeptides are standard techniques. Such techniques usually requires joining the sequences in a way so that they are in both in the same reading frame, and expression of the fused polypeptide under the control of the same promoter(s) and terminator.

30           Polypeptides of the present invention also include truncated forms of the neublastin polypeptide. In such truncated molecules, one or more amino acids have been deleted from the N-terminus or the C-terminus, preferably the N-terminus. Truncated neublastin polypeptides



include the human polypeptides designated herein as NBN112 (SEQ ID NO:35), NBN111 (SEQ ID NO:36), NBN110 (SEQ ID NO:37), NBN109 (SEQ ID NO:38), NBN108 (SEQ ID NO:39), NBN107 (SEQ ID NO:40), NBN106 (SEQ ID NO:41), NBN105 (SEQ ID NO:42), NBN104 (SEQ ID NO:43), NBN103 (SEQ ID NO:44), NBN102 (SEQ ID NO:45), NBN101 (SEQ ID NO:46), NBN100 (SEQ ID NO:47) and NBN99 (SEQ ID NO:48), and the corresponding homologs of these truncated human neublastin polypeptides, including but not limited to rat and mouse neublastin.

For example, the invention includes a truncated neublastin polypeptide whose amino terminus lacks one or more amino-terminal amino acids of a neublastin polypeptide. That is, the truncated neublastin polypeptide contains the seven cysteine domain of neublastin. In some embodiments, the truncated neublastin polypeptide includes an amino acid sequence with at least 70% homology to amino acids 12-113 of SEQ ID NO:12, *i.e.*, NBN102 (SEQ ID NO:45). Preferably, the truncated neublastin polypeptide is at least 85% homologous to amino acids 12-113 of SEQ ID NO:12. More preferably, the truncated neublastin polypeptide is at least 95% homologous to amino acids 12-113 of SEQ ID NO:12. Most preferably, the truncated neublastin polypeptide is at least 99% homologous to amino acids 12-113 of SEQ ID NO:12. Further similar examples of truncated neublastin include, *e.g.*, polypeptides that include amino acids 42-140 of SEQ ID NO:9, amino acids 113-224 of SEQ ID NO:16, amino acids 113-224 of SEQ ID NO:34. Additional specific examples include but are not limited to human NBN99 (SEQ ID NO:48), NBN100 (SEQ ID NO:47), NBN101 (SEQ ID NO:46), NBN102 (SEQ ID NO:45), NBN103 (SEQ ID NO:44), NBN104 (SEQ ID NO:43), NBN105 (SEQ ID NO:42), NBN106 (SEQ ID NO:41), NBN107 (SEQ ID NO:40), NBN108 (SEQ ID NO:39), NBN109 (SEQ ID NO:38), NBN110 (SEQ ID NO:37), NBN111 (SEQ ID NO:36) and NBN112 (SEQ ID NO:35), described above, or homologs or derivatives thereof. Truncated NBN polypeptides may be synthetic, expressed from cloned DNA constructs, or result from enzymatic cleavage of mature NBN polypeptides.

In preferred embodiments, the truncated neublastin polypeptide includes at least the 85 carboxy terminal amino acids of a neublastin polypeptide. In preferred embodiments, it includes at least the carboxy terminal 98, 99, 100, 101, 102, 103, 104, 105, 106, 107, 108, 109, 110, 111 or 112 amino acids of a neublastin polypeptide.

In preferred embodiments, the truncated neublastin polypeptide binds to a RET polypeptide, preferably where the RET polypeptide is expressed on the surface of a mammalian cell, such as a neuron.

The truncated neublastin can be prepared using recombinant expression of a nucleic acid encoding a truncated neublastin polypeptide using methods known in the art and sequences provided herein.

Alternatively, the truncated neublastin polypeptide may be obtained by providing a mature neublastin polypeptide, such as NBN113 of SEQ ID NO:12, and contacting the mature neublastin polypeptide with at least one protease under conditions sufficient to produce the truncated neublastin polypeptide. Preferably, at least one protease is an exoprotease, and contacting the neublastin polypeptide results in formation of an exopeptidase neublastin polypeptide digestion product that can be further digested with a dipeptidyl peptidase. In alternative embodiments, the protease is aminopeptidase, Endo Lys C or trypsin.

#### Amino Acid Sequence Homology

The degree to which a candidate polypeptide shares homology with a neublastin polypeptide of the invention is determined as the degree of identity between two amino acid sequences. A high level of sequence identity indicates a likelihood that the first sequence is derived from the second sequence.

Identity is determined by computer analysis, such as, without limitations, the ClustalX computer alignment program (*Thompson JD, Gibson TJ, Plewniak F, Jeanmougin F, & Higgins DG: The ClustalX windows interface: flexible strategies for multiple sequence alignment aided by quality analysis tools; Nucleic Acids Res.* 1997, 25 (24): 4876-82), and the default parameters suggested herein. Using this program, the mature part of a polypeptide encoded by an analogous DNA sequence of the invention exhibits a degree of identity of at least 90%, more preferably at least 95%, more preferably at least 98%, most preferably at least 99% with the amino acid sequence presented herein as SEQ ID NO:2, SEQ ID NO:4; SEQ ID NO:5; SEQ ID NO:6; SEQ ID NO:7; SEQ ID NO:9; SEQ ID NO:10; SEQ ID NO:11; SEQ ID NO:12, SEQ ID NO:16, SEQ ID NO:34, SEQ ID NO:35, SEQ ID NO:36, SEQ ID NO:37, SEQ ID NO:38, ID NO:39; SEQ ID NO:40; SEQ ID NO:41; SEQ ID NO:42, SEQ ID NO:43, SEQ ID NO:44; SEQ ID NO:45; SEQ ID NO:46; SEQ ID NO:47 or SEQ ID NO:48.

Based on the homology determination it has been confirmed that the neublastin polypeptide of the invention, belonging to the TGF- $\beta$  superfamily, is related to the GDNF subfamily, but represents a distinct member of this subfamily.

#### **Bioactive Polypeptides**

5        The polypeptide of the invention may be provided on any bioactive form, including the form of pre-pro-proteins, pro-proteins, mature proteins, glycosylated proteins, non-glycosylated proteins, phosphorylated proteins, non-phosphorylated proteins, truncated forms, or any other posttranslational modified protein. A bioactive neublastin polypeptide includes a polypeptide that, when dimerized, alone or in the presence of a cofactor (such as GFR $\alpha$ 3, or RET), binds to  
10    RET, induces dimerization of RET, and autophosphorylation of RET.

The polypeptide of the invention may in particular be a N-glycosylated polypeptide, which polypeptide preferably is glycosylated at the N-residues indicated in the sequence listings.

In a preferred embodiment, the polypeptide of the invention has the amino acid  
15    sequence presented as SEQ ID NO:9, holding a glycosylated asparagine residue at position 122; the amino acid sequence presented as SEQ ID NO:10, holding a glycosylated asparagine residue at position 122; the amino acid sequence presented as SEQ ID NO:11, holding a glycosylated asparagine residue at position 98; or the amino acid sequence presented as SEQ ID NO:12, holding a glycosylated asparagine residue at position 95.

20        This invention also contemplates neublastin fusion proteins, such as Ig-fusions, as described, *e.g.*, in United States patent 5,434,131, herein incorporated by reference, or serum albumin fusions.

In one embodiment, the invention provides a polypeptide having the amino acid sequence shown as SEQ ID NO:2, or an amino acid sequence which is at least about 85%, preferably at  
25    least about 90%, more preferably at least about 95%, more preferably at least about 98%, and most preferably at least about 99% homologous to the sequence presented as SEQ ID NO:2.

In another embodiment, the invention provides a polypeptide having the amino acid sequence of SEQ ID NO:4, or an amino acid sequence which is at least about 85%, preferably at least about 90%, more preferably at least about 95%, more preferably at least about 98%, and  
30    most preferably at least about 99% homologous to the sequence presented as SEQ ID NO:4.

In a third embodiment, the invention provides a polypeptide having the amino acid sequence of SEQ ID NO:5, or an amino acid sequence which is at least about 85%, preferably at

least about 90%, more preferably at least about 95%, more preferably at least about 98%, and most preferably at least about 99% homologous to the sequence presented as SEQ ID NO:5.

In a fourth embodiment, the invention provides a polypeptides having the amino acid sequence of SEQ ID NO:6, or an amino acid sequence which is at least about 85%, preferably  
5 at least about 90%, more preferably at least about 95%, more preferably at least about 98%, and most preferably at least about 99% homologous to the sequence presented as SEQ ID NO:6.

In a fifth embodiment, the invention provides a polypeptides having the amino acid sequence of SEQ ID NO:7, or an amino acid sequence which is at least about 85%, preferably at least about 90%, more preferably at least about 95%, more preferably at least about 98%, and  
10 most preferably at least about 99% homologous to the sequence presented as SEQ ID NO:7.

The neublastin polypeptide of the invention includes allelic variants, e.g., the polypeptide amino acid sequences of SEQ ID NOS. 5-7, in which the first Xaa designates Asn or Thr, and the second Xaa designates Ala or Pro.

In a sixth embodiment, the invention provides a polypeptides having the amino acid  
15 sequence of SEQ ID NO:9, or an amino acid sequence which is at least about 85%, preferably at least about 90%, more preferably at least about 95%, more preferably at least about 98%, and most preferably at least about 99% homologous to the sequence presented as SEQ ID NO:9.

In a seventh embodiment, the invention provides a polypeptide having the amino acid sequence of SEQ ID NO:10, or an amino acid sequence at least about 85%, preferably at least  
20 about 90%, more preferably at least about 95%, more preferably at least about 98%, and most preferably at least about 99% homologous to the sequence presented as SEQ ID NO:10.

In a eighth embodiment, the invention provides a polypeptide having the amino acid sequence of SEQ ID NO:11, or an amino acid sequence at least about 85%, preferably at least  
25 about 90%, more preferably at least about 95%, more preferably at least about 98%, and most preferably at least about 99% homologous to the sequence presented as SEQ ID NO:11.

In a ninth embodiment, the invention provides a polypeptide having the amino acid sequence of SEQ ID NO:12, or an amino acid sequence at least about 85%, preferably at least  
about 90%, more preferably at least about 95%, more preferably at least about 98%, and most preferably at least about 99% homologous to the sequence presented as SEQ ID NO:12.

30 In a tenth embodiment, the invention provides a polypeptide having the amino acid sequence of SEQ ID NO:16, or an amino acid sequence at least about 85%, preferably at least about 90%, more preferably at least about 95%, more preferably at least about 98%, and most

preferably at least about 99% homologous to the sequence presented as SEQ ID NO:16, which is a pre-pro-neublastin of murine origin.

In further embodiments, the invention provides a polypeptide having the amino acid sequence of any one of SEQ ID NOS:35-48, or an amino acid sequence at least about 85%,  
5 preferably at least about 90%, more preferably at least about 95%, more preferably at least about 98%, and most preferably at least about 99% homologous to any one of the sequences presented as SEQ ID NOS:35-48.

In another embodiment, the polypeptide of the invention holds the GDNF subfamily fingerprint, *i.e.* the amino acid residues underlined in Table 3, shown in the Examples.

10 In a further embodiment, the invention provides a polypeptide encoded by a polynucleotide sequence capable of hybridizing under high stringency conditions with the polynucleotide sequence presented as SEQ ID NO:1, its complementary strand, or a sub-sequence thereof. In a preferred embodiment, the polypeptide of the invention is encoded by a polynucleotide sequence being at least 70% homologous to the polynucleotide sequence  
15 presented as SEQ ID NO:1. In its most preferred embodiment, the polypeptide of the invention is encoded by the polynucleotide sequence presented as SEQ ID NO:1.

In a yet further embodiment, the invention provides novel polypeptides encoded by a polynucleotide sequence capable of hybridizing under high stringency conditions with the polynucleotide sequence presented as SEQ ID NO:3, its complementary strand, or a sub-  
20 sequence thereof. In a preferred embodiment, the polypeptide of the invention is encoded by a polynucleotide sequence being at least 70% homologous to the polynucleotide sequence presented as SEQ ID NO:3. In its most preferred embodiment, the polypeptide of the invention is encoded by the polynucleotide sequence presented as SEQ ID NO:3.

In a more preferred embodiment, the polypeptide of the invention includes the amino  
25 acid sequence of a mature Neublastin polypeptide. Even more preferably, the invention includes the amino acid sequence of a truncated form of the Neublastin polypeptide that includes the seven conserved cysteine residues present in the amino acid sequence of Neublastin proteins.

In a still further embodiment, the invention provides novel polypeptides encoded by a polynucleotide sequence capable of hybridizing under high stringency conditions with the  
30 polynucleotide sequence presented as SEQ ID NO:8, its complementary strand, or a sub-sequence thereof. In a preferred embodiment, the polypeptide of the invention is encoded by a polynucleotide sequence being at least 70% homologous to the polynucleotide sequence

presented as SEQ ID NO:8. In its most preferred embodiment, the polypeptide of the invention is encoded by the polynucleotide sequence presented as SEQ ID NO:8.

In a still further embodiment, the invention provides novel polypeptides encoded by a polynucleotide sequence capable of hybridizing under high stringency conditions with the polynucleotide sequence presented as SEQ ID NO:15, its complementary strand, or a sub-  
5 sequence thereof. In a preferred embodiment, the polypeptide of the invention is encoded by a polynucleotide sequence being at least 70% homologous to the polynucleotide sequence presented as SEQ ID NO:15. In its most preferred embodiment, the polypeptide of the invention is encoded by the polynucleotide sequence presented as SEQ ID NO:15.

10

### **Biological Origin**

The polypeptide of the invention may be isolated from mammalian cells, preferably from a human cell or from a cell of murine origin.

In a most preferred embodiment, the polypeptide of the invention may be isolated from  
15 human heart tissue, from human skeletal muscle, from human pancreas, or from human brain tissue, in particular from caudate nucleus or from thalamus, or it may be obtained from DNA of mammalian origin, as discussed in more detail below.

### **Neurotrophic Activity**

Neublastin polypeptides, including truncated neublastin polypeptides, of the invention are useful for moderating metabolism, growth, differentiation, or survival of a nerve or neuronal cell. In particular, neublastin polypeptides are used to treating or to alleviate a disorder or disease of a living animal, *e.g.*, a human, which disorder or disease is responsive to the activity of a neurotrophic agents. Such treatments and methods are described in more details below.

25

### **Antibodies**

Neublastin polypeptides or polypeptide fragments of the invention are used to produce neublastin-specific antibodies. As used herein, a "neublastin-specific antibody" is an antibody, *e.g.*, a polyclonal antibody or a monoclonal antibody, that is immunoreactive to a neublastin  
30 polypeptide or polypeptide fragment, or that binds with specificity to an epitope of a neublastin polypeptide.

The preparation of polyclonal and monoclonal antibodies is well known in the art. Polyclonal antibodies may in particular be obtained as described by, *e.g.*, *Green et al.*,: "Production of Polyclonal Antisera" in *Immunochemical Protocols* (*Manson*, Ed.); Humana Press, 1992, pages 1-5; by *Coligan et al.*,: "Production of Polyclonal Antisera in Rabbits, Rats, Mice and Hamsters" in *Current Protocols in Immunology*, 1992, Section 2.4.1, and by Ed Harlow and David Lane (Eds.) in "Antibodies; A laboratory manual" Cold Spring Harbor Lab. Press 1988. These protocols are hereby incorporated by reference. Monoclonal antibodies may in particular be obtained as described by, *e.g.*, *Kohler & Milstein*, *Nature*, 1975, 256:495; *Coligan et al.*, in *Current Protocols in Immunology*, 1992, Sections 2.5.1 - 2.6.7; and *Harlow et al.*, in *Antibodies: A Laboratory Manual*; Cold Spring Harbor, Pub., 1988, page 726; which protocols are hereby incorporated by reference.

Briefly, monoclonal antibodies may be obtained by injecting, *e.g.*, mice with a composition comprising an antigen, verifying the presence of antibody production by removing a serum sample, removing the spleen to obtain B lymphocytes, fusing the B lymphocytes with myeloma cells to produce hybridomas, cloning the hybridomas, selecting positive clones that produce the antibodies to the antigen, and isolating the antibodies from the hybridoma cultures.

Monoclonal antibodies can be isolated and purified from hybridoma cultures by a variety of well-established techniques, including affinity chromatography with protein A Sepharose, size-exclusion chromatography, and ion-exchange chromatography, see. *e.g.* *Coligan et al.* in *Current Protocols in Immunology*, 1992, Sections 2.7.1 - 2.7.12, and Sections 2.9.1 - 2.9.3; and *Barnes et al.*: "Purification of Immunoglobulin G (IgG)" in *Methods in Molecular Biology*; Humana Press, 1992, Vol. 10, Pages 79-104. Polyclonal or monoclonal antibodies may optionally be further purified, *e.g.* by binding to and elution from a matrix to which is bound the polypeptide against which the antibodies were raised.

Antibodies which bind to the neublastin polypeptide of the invention can be prepared using an intact polypeptide or fragments containing small peptides of interest as the immunising antigen. The polypeptide used to immunise an animal may be obtained by recombinant DNA techniques or by chemical synthesis, and may optionally be conjugated to a carrier protein. Commonly used carrier proteins which are chemically coupled to the peptide include keyhole limpet hemocyanin (KLH), thyroglobulin, bovine serum albumin (BSA), and tetanus toxoid. The coupled peptide may then be used to immunise the animal, which may in particular be a mouse, a rat, a hamster or a rabbit.

In one embodiment, antibodies are produced using the following peptides: Peptide 1: CRPTRYEAVSFMDVNST (amino acids 108-124 of SEQ ID NO:9); or Peptide 2: ALRPPGSRPVSQPC (amino acids 93-107 of SEQ ID NO:9). Methods for producing antibodies using these polypeptides are described in Example 10.

- 5 Rabbit polyclonal antibodies were also generated to the following peptides:  
 Peptide R27: GPGSRARAAGARGC (amino acids 30-43 of SEQ ID NO:9);  
 Peptide R28: LGHRSEDELVRFRFC (amino acids 57-70 of SEQ ID NO:9);  
 Peptide R29: CRRARSPHDLSL (amino acids 74-85 of SEQ ID NO:9);  
 Peptide R30: LRPPGSRPVSQPC (amino acids 94-107 of SEQ ID NO:9); and  
 10 Peptide R31: STWRTVDRLSATAC (amino acids 123-136 of SEQ ID NO:9).

Of this group, only peptides R30 and R31, relatively close to the C-terminus, recognized the denatured protein under reducing conditions on a Western blot.

- Additional neublastin-derived peptides also were derived from the NBN protein, as detailed below, which are predicted surface exposed loops based on the known GDNF structure  
 15 (Eigenbrot and Gerber, Nat. Struct. Biol., 4, pp. 435-438 (1997)), and are thus useful for antibody generation:

Region 1: CRLRSQLVVPVRALGLGHRSEDELVRFRFC (AA43-70 of SEQ ID NO:9)  
 Region 2: CRRARSPHDLSLASLLGAGALRPPGSRPVSQPC (AA74-107 of SEQ ID NO:9)  
 Region 3: CRPTRYEAVSFMDVNSTWRTVDRLSATAC (AA108-136 of SEQ ID NO:9)

20

- In another aspect of the invention, antibodies which specifically bind neublastin or neublastin-derived peptides may be used for detecting the presence of such neublastin neurotrophic factors in various media, and in particular for the diagnosis of conditions or diseases associated with the neublastin molecules of the invention. A variety of protocols for  
 25 such detection, including ELISA, RIA and FACS, are known in the art.

The antibodies of this invention may also be used for blocking the effect of the neublastin neurotrophic factor, and may in particular be neutralizing antibodies.

#### **Methods of Producing the Polypeptides of the Invention**

- 30 A cell comprising a DNA sequence encoding a neublastin polypeptide of the invention is cultured under conditions permitting the production of the polypeptide, followed by recovery of the polypeptide from the culture medium, as detailed below. When cells are to be



genetically modified for the purposes of producing a neublastin polypeptide, the cells may be modified by conventional methods or by gene activation.

According to conventional methods, a DNA molecule that contains a neublastin cDNA or genomic DNA sequence may be contained within an expression construct and transfected  
5 into cells by standard methods including, but not limited to, liposome-, polybrene-, or DEAE dextran-mediated transfection, electroporation, calcium phosphate precipitation, microinjection, or velocity driven microprojectiles ("biolistics"). Alternatively, one could use a system that delivers DNA by viral vector. Viruses known to be useful for gene transfer include adenoviruses, adeno-associated virus, lentivirus, herpes virus, mumps virus, poliovirus,  
10 retroviruses, Sindbis virus, and vaccinia virus such as canary pox virus, as well as Baculovirus infection of insect cells, in particular Sf9 insect cells.

Alternatively, the cells may be modified using a gene activation ("GA") approach, such as described in United States patents 5,733,761 and 5,750,376, each incorporated herein by reference.

15 Accordingly, the term "genetically modified," as used herein in reference to cells, is meant to encompass cells that express a particular gene product following introduction of a DNA molecule encoding the gene product and/or regulatory elements that control expression of a coding sequence for the gene product. The DNA molecule may be introduced by gene targeting, allowing incorporation of the DNA molecule at a particular genomic site.

20

#### **Recombinant Expression Vectors**

In a further aspect the invention provides a recombinant expression vector comprising the polynucleotide of the invention. The recombinant expression vector of the invention may be any suitable eukaryotic expression vector. Preferred recombinant expression vectors are the  
25 ubiquitin promoter containing vector pTEJ-8 (FEBS Lett. 1990 **267** 289-294), and derivatives hereof, *e.g.* pUbi1Z. A preferred commercially available eukaryotic expression vector is *e.g.* the virus promoter containing vector pcDNA-3 (available from Invitrogen). Another preferred expression vector uses SV40 early and adenovirus major late promoters (derived from plasmid pAD2beta; Norton and Coffin, *Mol. Cell. Biol.* **5**: 281 (1985)).

30 This invention also provides prokaryotic expression vectors and synthetic genes (syngenes) with codon optimization for prokaryotic expression. Syngenes were constructed with lower GC content and preferred bacterial (*e.g.*, *E. coli*) codons. The syngene has been

cloned into two vectors, pET19b and pMJB164, a derivative of pET19b. The construction with pET19b is shown in FIG. 14. In this construct, the sequence encoding the NBN113 domain of neublastin is directly fused to an initiating methionine. The construction with pMJB164 is shown in FIG. 15.

5

### Production Cells

In a yet further aspect the invention provides a production cell genetically manipulated to comprise the isolated polynucleotide sequence of the invention, and/or or a recombinant expression vector of the invention. The cell of the invention may in particular be genetically manipulated to transiently or stably express, over-express or co-express polypeptide of the invention. Methods for generating transient and stable expression are known in the art.

The polynucleotide of the invention may be inserted into an expression vector, *e.g.* a plasmid, virus or other expression vehicle, and operatively linked to expression control sequences by ligation in a way that expression of the coding sequence is achieved under conditions compatible with the expression control sequences. Suitable expression control sequences include promoters, enhancers, transcription terminators, start codons, splicing signals for introns, and stop codons, all maintained in the correct reading frame of the polynucleotide of the invention so as to permit proper translation of mRNA. Expression control sequences may also include additional components such as leader sequences and fusion partner sequences.

The promoter may in particular be a constitutive or an inducible promoter. When cloning in bacterial systems, inducible promoters such as pL of bacteriophage  $\lambda$ , plac, ptrp, ptac (ptrp-lac hybrid promoter), may be used. When cloning in mammalian systems, promoters derived from the genome of mammalian cells, *e.g.* the ubiquitin promoter, the TK promoter, or the metallothionein promoter, or from mammalian viruses, *e.g.* the retrovirus long terminal repeat, the adenovirus late promoter or the vaccinia virus 7.5K promoter, may be used. Promoters obtained by recombinant DNA or synthetic techniques may also be used to provide for transcription of the polynucleotide of the invention.

Suitable expression vectors typically comprise an origin of expression, a promoter as well as specific genes which allow for phenotypic selection of the transformed cells, and include vectors like the T7-based expression vector for expression in bacteria (*Rosenberg et al*; Gene 1987 56 125), the pTEJ-8, pUbi1Z, pcDNA-3 and pMSXND expression vectors for

expression in mammalian cells (*Lee and Nathans, J. Biol. Chem.* 1988 **263** 3521), baculovirus derived vectors for expression in insect cells, and the oocyte expression vector PTLN (*Lorenz C, Pusch M & Jentsch T J: Heteromultimeric CLC chloride channels with novel properties; Proc. Natl. Acad. Sci. USA* 1996 **93** 13362-13366).

5 In a preferred embodiment, the cell of the invention is a eukaryotic cell, *e.g.*, a mammalian cell, *e.g.*, a human cell, an oocyte, or a yeast cell. The cell of the invention may be without limitation a human embryonic kidney (HEK) cell, *e.g.*, a HEK 293 cell, a BHK21 cell, a Chinese hamster ovary (CHO) cell, a *Xenopus laevis* oocyte (XLO) cell. In one embodiment, the cell of the invention is a fungal cell, *e.g.*, a filamentous fungal cell. In yet another  
10 embodiment, the cell is an insect cell, most preferably the Sf9 cell. Additional mammalian cells of the invention are PC12, HiB5, RN33b cell lines, human neural progenitor cells, and other cells derived from human cells, especially neural cells.

Examples of primary or secondary cells include fibroblasts, epithelial cells including mammary and intestinal epithelial cells, endothelial cells, formed elements of the blood  
15 including lymphocytes and bone marrow cells, glial cells, hepatocytes, keratinocytes, muscle cells, neural cells, or the precursors of these cell types. Examples of immortalized human cell lines useful in the present methods include, but are not limited to, Bowes Melanoma cells (ATCC Accession No. CRL 9607), Daudi cells (ATCC Accession No. CCL 213), HeLa cells and derivatives of HeLa cells (ATCC Accession Nos. CCL 2, CCL 2.1, and CCL 2.2), HL-60  
20 cells (ATCC Accession No. CCL 240), HT-1080 cells (ATCC Accession No. CCL 121), Jurkat cells (ATCC Accession No. TIB 152), KB carcinoma cells (ATCC Accession No. CCL 17), K-562 leukemia cells (ATCC Accession No. CCL 243), MCF-7 breast cancer cells (ATCC Accession No. BTH 22), MOLT-4 cells (ATCC Accession No. 1582), Namalwa cells (ATCC Accession No. CRL 1432), Raji cells (ATCC Accession No. CCL 86), RPMI 8226  
25 cells (ATCC Accession No. CCL 155), U-937 cells (ATCC Accession No. CRL 1593), WI-38VA13 sub line 2R4 cells (ATCC Accession No. CLL 75.1), and 2780AD ovarian carcinoma cells (Van der Blick *et al.*, *Cancer Res.* 48: 5927-5932, 1988), as well as heterohybridoma cells produced by fusion of human cells and cells of another species. Secondary human fibroblast strains, such as WI-38 (ATCC Accession No. CCL 75) and  
30 MRC-5 (ATCC Accession No. CCL 171), may also be used.

When the cell of the invention is an eukaryotic cell, incorporation of the heterologous polynucleotide of the invention may in particular be carried out by infection (employing a virus

vector), by transfection (employing a plasmid vector), using calcium phosphate precipitation, microinjection, electroporation, lipofection, or other physical-chemical methods known in the art.

In a more preferred embodiment the isolated polynucleotide sequence of the invention, and/or or a recombinant expression vector of the invention are transfected in a mammalian host cell, a neural progenitor cell, an astrocyte cell, a T-cell, a hematopoietic stem cell, a non-dividing  
5 cell, or a cerebral endothelial cell, comprising at least one DNA molecule capable of mediating cellular immortalization and/or transformation.

Activation of an endogenous gene in a host cell may be accomplished by the introducing regulatory elements, in particular by the introducing a promoter capable of effecting transcription  
10 of an endogenous gene encoding the neublastin polypeptide of the invention.

#### **Pharmaceutical Compositions**

In another aspect the invention provides novel pharmaceutical compositions comprising a therapeutically effective amount of the polypeptide of the invention.

15 For use in therapy the polypeptide of the invention may be administered in any convenient form. In a preferred embodiment, the polypeptide of the invention is incorporated into a pharmaceutical composition together with one or more adjuvants, excipients, carriers and/or diluents, and the pharmaceutical composition prepared by the skilled person using conventional methods known in the art.

20 Such pharmaceutical compositions may comprise the polypeptide of the invention, or antibodies hereof. The composition may be administered alone or in combination with at one or more other agents, drugs or hormones.

The pharmaceutical composition of this invention may be administered by any suitable route, including, but not limited to oral, intravenous, intramuscular, inter-arterial,  
25 intramedullary, intrathecal, intraventricular, transdermal, subcutaneous, intraperitoneal, intranasal, arterial, topical, sublingual or rectal application, buccal, vaginal, intraorbital, intracerebral, intracranial, intraspinal, intraventricular, intracisternal, intracapsular, intrapulmonary, transmucosal, or via inhalation.

Intrapulmonary delivery methods, apparatus and drug preparation are described, for  
30 example, in U.S. Patent Nos. 5, 785, 049, 5,780,019, and 5,775,320, each incorporated herein by reference. Administration may be by periodic injections of a bolus of the preparation, or may be made more continuous by intravenous or intraperitoneal administration from a reservoir

which is external (*e.g.*, an IV bag) or internal (*e.g.*, a bioerodable implant, a bioartificial organ, or a colony of implanted neublastin production cells). See, *e.g.*, U.S. Patents 4,407,957, 5,798,113, and 5,800,828, each incorporated herein by reference. Intrapulmonary delivery methods and apparatus are described, for example, in U.S. Patents 5,654,007, 5,780,014, and 5,814,607, each incorporated herein by reference.

In particular, administration of a neublastin according to this invention may be achieved using any suitable delivery means, including:

- (a) pump (see, *e.g.*, Annals of Pharmacotherapy, 27:912 (1993); Cancer, 41:1270 (1993); Cancer Research, 44:1698 (1984), incorporated herein by reference),
- 10 (b), microencapsulation (see, *e.g.*, United States patents 4,352,883; 4,353,888; and 5,084,350, herein incorporated by reference),
- (c) continuous release polymer implants (see, *e.g.*, Sabel, United States patent 4,883,666, incorporated herein by reference),
- (d) macroencapsulation (see, *e.g.*, United States patents 5,284,761, 5,158,881,
- 15 4,976,859 and 4,968,733 and published PCT patent applications WO92/19195, WO 95/05452, each incorporated herein by reference);
- (e) naked or unencapsulated cell grafts to the CNS (see, *e.g.*, United States patents 5,082,670 and 5,618,531, each incorporated herein by reference); or
- (f) injection, either subcutaneously, intravenously, intra-arterially, intramuscularly, or
- 20 to other suitable site;
- (g) oral administration, in capsule, liquid, tablet, pill, or prolonged release formulation.

In one embodiment of this invention, a neublastin polypeptide is delivered directly into the CNS, preferably to the brain ventricles, brain parenchyma, the intrathecal space or other suitable CNS location, most preferably intrathecally.

25 In another preferred embodiment, the present neublastin polypeptide is given by systemic delivery via intramuscular injection, subcutaneous injection, intravenous injection, or intravenous infusion.

Other useful parenteral delivery systems include ethylene-vinyl acetate copolymer particles, osmotic pumps, implantable infusion systems, pump delivery, encapsulated cell

30 delivery, liposomal delivery, needle-delivered injection, needle-less injection, nebulizer, aerosolizer, electroporation, and transdermal patch.

Further details on techniques for formulation and administration may be found in the latest edition of Remington's Pharmaceutical Sciences (Maack Publishing Co., Easton, PA).

The active ingredient may be administered in one or several doses per day. Currently contemplated appropriate dosages are between 0.5 ng neublastin/kg body weight to about 50  
5  $\mu\text{g/kg}$  per administration, and from about 1.0 ng/kg to about 100  $\mu\text{g/kg}$  daily. When delivered directly to the CNS, the neublastin pharmaceutical composition should provide a local concentration of neurotrophic factor of from about 5 ng/ml cerebrospinal fluid ("CSF") to 25 ng/ml CSF.

The dose administered must of course be carefully adjusted to the age, weight and  
10 condition of the individual being treated, as well as the route of administration, dosage form and regimen, and the result desired, and the exact dosage should of course be determined by the practitioner.

In further embodiments, the neublastin polypeptide of the invention may be administered by genetic delivery, using cell lines and vectors as described below under  
15 methods of treatment. To generate such therapeutic cell lines, the polynucleotide of the invention may be inserted into an expression vector, *e.g.* a plasmid, virus or other expression vehicle, and operatively linked to expression control sequences by ligation in a way that expression of the coding sequence is achieved under conditions compatible with the expression control sequences. Suitable expression control sequences include promoters, enhancers,  
20 transcription terminators, start codons, splicing signals for introns, and stop codons, all maintained in the correct reading frame of the polynucleotide of the invention so as to permit proper translation of mRNA. Expression control sequences may also include additional components such as leader sequences and fusion partner sequences.

The promoter may in particular be a constitutive or an inducible promoter. Constitutive  
25 promoters could be synthetic, viral or derived from the genome of mammalian cells, *e.g.* the human ubiquitin promoter. In a preferred embodiment the therapeutic cell line will be a human immortalised neural cell line expressing the polypeptide of the invention. For implantation, we contemplate implanting between about  $10^5$  to  $10^{10}$  cells, more preferably  $10^6$  to about  $10^8$  cells.

### 30 Methods of Treatment

The present invention, which relates to polynucleotides and proteins, polypeptides, peptide fragments or derivatives produced therefrom, as well as to antibodies directed against

such proteins, peptides or derivatives, may be used for treating or alleviating a disorder or disease of a living animal body, including a human, which disorder or disease is responsive to the activity of neurotrophic agents.

The polypeptides of the present invention may be used directly via, *e.g.*, injected,  
5 implanted or ingested pharmaceutical compositions to treat a pathological process responsive to the neublastin polypeptides.

The polynucleotide of the invention, including the complementary sequences thereof, may be used for the expression of the neurotrophic factor of the invention. This may be achieved by cell lines expressing such proteins, peptides or derivatives of the invention, or by  
10 virus vectors encoding such proteins, peptides or derivatives of the invention, or by host cells expressing such proteins, peptides or derivatives. These cells, vectors and compositions may be administered to treatment target areas to affect a disease or disorder process responsive to the neublastin polypeptides.

Suitable expression vectors may be derived from lentiviruses, retroviruses,  
15 adenoviruses, herpes or vaccinia viruses, or from various bacterially produced plasmids may be used for *in vivo* delivery of nucleotide sequences to a whole organism or a target organ, tissue or cell population. Other methods include, but are not limited to, liposome transfection, electroporation, transfection with carrier peptides containing nuclear or other localizing signals, and gene delivery via slow-release systems. In still another aspect of the invention, "antisense"  
20 nucleotide sequences complementary to the neublastin gene or portions thereof, may be used to inhibit or enhance neublastin expression.

In yet another aspect the invention relates to a method of treating or alleviating a disorder or disease of a living animal body, including a human, which disorder or disease is responsive to the activity of neurotrophic agents.

25 The disorder or disease may in particular be damage of the nervous system caused by trauma, surgery, ischemia, infection, reperfusion, metabolic disease, nutritional deficiency, malignancy or a toxic agent, or a genetic or idiopathic processes.

The damage may in particular have occurred to sensory neurons or retinal ganglion cells, including neurons in the dorsal root ganglion or in any of the following tissues: the  
30 geniculate, petrosal and nodose ganglia; the vestibuloacoustic complex of the VIIIth cranial nerve; the ventrolateral pole of the maxillomandibular lobe of the trigeminal ganglion; and the mesencephalic trigeminal nucleus.

In a preferred embodiment of the method of the invention, the disease or disorder is a neurodegenerative disease involving lesioned or traumatized neurons, such as traumatic lesions of peripheral nerves, the medulla, and/or the spinal cord, cerebral ischemic neuronal damage, neuropathy and especially peripheral neuropathy, peripheral nerve trauma or injury, ischemic stroke, acute brain injury, acute spinal cord injury, nervous system tumors, multiple sclerosis, exposure to neurotoxins, metabolic diseases such as diabetes or renal dysfunctions and damage caused by infectious agents, neurodegenerative disorders including Alzheimer's disease, Huntington's disease, Parkinson's disease, Parkinson-Plus syndromes, progressive Supranuclear Palsy (Steele-Richardson-Olszewski Syndrome), Olivopontocerebellar Atrophy (OPCA), Shy-Drager Syndrome (multiple systems atrophy), Guamanian parkinsonism dementia complex, amyotrophic lateral sclerosis, or any other congenital or neurodegenerative disease, and memory impairment connected to dementia.

In a preferred embodiment, treatment is contemplated of sensory and/or autonomic system neurons. In another preferred embodiment, treatment is contemplated of motor neuron diseases such as amyotrophic lateral sclerosis ("ALS") and spinal muscular atrophy. In yet another preferred embodiment, use is contemplated of the neublastin molecules of this invention to enhance nerve recovery following traumatic injury. In one embodiment use is contemplated of a nerve guidance channel with a matrix containing neublastin polypeptides. Such nerve guidance channels are disclosed, *e.g.*, United States Patent No. 5,834,029, incorporated herein by reference.

In a preferred embodiment, the polypeptides and nucleic acids of this invention (and pharmaceutical compositions containing same) are used in the treatment of peripheral neuropathies. Among the peripheral neuropathies contemplated for treatment with the molecules of this invention are trauma-induced neuropathies, *e.g.*, those caused by physical injury or disease state, physical damage to the brain, physical damage to the spinal cord, stroke associated with brain damage, and neurological disorders related to neurodegeneration.

Treatment also is contemplated of chemotherapy-induced neuropathies (such as those caused by delivery of chemotherapeutic agents, *e.g.*, taxol or cisplatin); toxin-induced neuropathies, drug-induced neuropathies, pathogen-induced (*e.g.*, virus induced) neuropathies, vitamin-deficiency-induced neuropathies; idiopathic neuropathies; and diabetic neuropathies. See, *e.g.*, United States Patent Nos. 5,496,804 and 5,916,555, each herein incorporated by reference.



Treatment also is contemplated of mono-neuropathies, mono-multiplex neuropathies, and poly-neuropathies, including axonal and demyelinating neuropathies, using the neublastin nucleotides and polypeptides of this invention.

In another preferred embodiment, the polypeptides and nucleic acids of this invention  
5 (and pharmaceutical compositions containing same) are used in the treatment of various disorders in the eye, including photoreceptor loss in the retina in patients afflicted with macular degeneration, retinitis pigmentosa, glaucoma, and similar diseases.

The present invention additionally provides a method for the prevention of the degenerative changes connected with the above diseases and disorders, by implanting into  
10 mammalian brain human vectors or cells capable of producing a biologically active form of neublastin or a precursor of neublastin, *i.e.* a molecule that can readily be converted to a biologically active form of neublastin by the body, or additionally cells that secrete neublastin may be encapsulated, *e.g.* into semipermeable membranes.

Suitable cells, including cells engineered to produce neublastin, can be grown *in vitro*  
15 for use in transplantation or engraftment into mammalian brain including human.

In a preferred embodiment, the gene encoding the polypeptide of the invention is transfected into a suitable cell line, *e.g.* into an immortalised rat neural stem cell line like HiB5 and RN33b, or into a human immortalised neural progenitor cell line, and the resulting cell line is implanted in the brain of a living body, including a human, to secrete the therapeutic  
20 polypeptide of the invention in the CNS, *e.g.* using the expression vectors described in International Patent Application WO 98/32869.

#### Methods of Diagnosis and Screening

A neublastin nucleic acid can be used to determine whether an individual is predisposed  
25 to developing a neurological disorder resulting from a defect in the neublastin gene, *e.g.*, an defect in a neublastin allele, which has been acquired by, *e.g.*, genetic inheritance, by abnormal embryonic development, or by acquired DNA damage. The analysis can be by, *e.g.*, detecting a deletion(s) or a point-mutation(s) within the neublastin gene, or by detecting the inheritance of such predisposition of such genetic defects with specific restriction fragment length  
30 polymorphisms (RFLPs), by detecting the presence or absence of a normal neublastin gene by hybridizing a nucleic acid sample from the patient with a nucleic acid probe(s) specific for the neublastin gene, and determining the ability of the probe to hybridize to the nucleic acid.

In particular, a neublastin nucleic acid can be used as a hybridization probe. Such hybridization assays may be used to detect, prognose, diagnose, or monitor the various conditions, disorders, or disease states associated with aberrant levels of the mRNAs encoding the Neublastin protein. A neublastin nucleic acid can be construed as a "marker" for neublastin neurotrophic factor-dependant physiological processes. These processes include, but are not limited to, "normal" physiological processes (*e.g.*, neuronal function) and pathological processes (*e.g.*, neurodegenerative disease). The characterization of a particular patient sub-population(s) with aberrant (*i.e.*, elevated or deficient) levels of the neublastin protein and or neublastin-encoding mRNA may lead to new disease classifications. By "aberrant levels," as defined herein, is meant an increased or decreased level relative to that in a control sample or individual not having the disorder determined by quantitative or qualitative means.

The neublastin nucleic acids and polypeptides of this invention may also be used to screen for and identify neublastin analogs, including small molecule mimetics of neublastin. In one contemplated embodiment, the invention provides a method for identifying a candidate compound that induces a neublastin-mediated biological effect, the method comprising the steps of providing a test cell which when contacted with neublastin is induced to express a detectable product, exposing the cell to the candidate compound, and detecting the detectable product. The expression of the detectable product is indicative of the ability of the candidate compound to induce the neublastin-mediated biological effect.

Further, the neublastin nucleic acids and polypeptides of this invention may be used on DNA chip or protein chips, or in computer programs to identify related novel gene sequences and proteins encoded by them, including allelic variants and single nucleotide polymorphisms ("SNPs"). Such methods are described, *e.g.*, in United States Patent Nos. 5,795,716; 5,754,524; 5,733,729; 5,800,992; 5,445,934; 5,525,464, each herein incorporated by reference.

## EXAMPLES

### Example 1: Methods for Isolating Neublastin Nucleic Acids

#### **Method 1: Rapid-Screening of Human Fetal Brain cDNA for the neublastin Gene**

A 290 bp fragment was identified in two high throughput genomic sequences (HTGS) submitted to GenBank (Accession No. AC005038 and AC005051) by its homology to human persephin. From the nucleic acid sequence of the 290 bp fragment, two neublastin specific

primers were synthesized. The neublastin top strand primer ("NBNint.sense") had the sequence 5'-CCT GGC CAG CCT ACT GGG-3' (SEQ ID NO:17). The neublastin bottom strand primer ("NBNint.antisense") had the sequence 5'-AAG GAG ACC GCT TCG TAG CG-3' (SEQ ID NO:18). With these primers, 96-well PCR reactions were performed.

5 A 96-well master plate, containing plasmid DNA from 500,000 cDNA clones, was loaded with approximately 5000 clones per well. A 96-well sub-plate was utilized with *E. coli* DH10B glycerol stock containing 50 clones per well.

A neublastin nucleic acid was identified by three rounds of amplification using polymerase chain reaction ("PCR") techniques; amplification increases the number of copies of  
10 the nucleic acid in the sample.

**Master Plate Screening:** Using the 96-well PCR screening technique described above, a human fetal brain cDNA master plate was screened with the gene-specific primers to isolate the human neublastin cDNA.

Thirty nanograms (30 ng) of human fetal brain cDNA (6 ng/μl; Origene Technologies)  
15 was obtained from the corresponding well of the master plate and placed in a total volume of 25 μl which contained the following reagents: 0.2 mM of each of the two aforementioned gene-specific primers (*i.e.*, NBNint.sense and NBNint.antisense), 1x standard PCR buffer (Buffer V, Advanced Biotechnologies, UK), 0.2 mM dNTPs (Amersham-Pharmacia), 0.1 M GC-Melt (Clontech Laboratories, USA); and 0.5 units of *Taq* DNA polymerase (5 U/μl; Advanced  
20 Biotechnologies, UK).

PCR thermocycling reactions were performed using the following procedure and conditions. DNA was initially denatured at 94°C for 3 minutes, and then followed by 35 cycles of denaturation at 94°C for 1 minute each, annealing at 55°C for 1 minute, a first extension at 72°C for 90 seconds; and a final extension at 72°C for 5 minutes. The products of 96  
25 individual PCR reactions were analysed by gel electrophoresis using a 2% agarose gel containing ethidium bromide stain. The 102 bp, positive PCR product seen from a well was found to correspond to a unique 96-well sub-plate.

The 102 bp nucleic acid fragment had the following sequence (SEQ ID NO:13):  
5' - CCTGGCCAGCCTACTGGGCGCCGGGGCCCTGCGACCGCCCCGGGCTCCCGGCCGTCAG  
30 CCAGCCCTGCTGCCGACCCACGCGCTACGAAGCGGTCTCCTT - 3'

**Sub-Plate Screening:** The 96-well human fetal brain sub-plate was screened by PCR-mediated amplification by placing 1 µl of the glycerol stock from the corresponding sub-plate well in a total volume of 25 µl which contained: 0.2 mM of each of the two gene-specific primers; 1x standard PCR buffer (Buffer V; Advanced Biotechnologies, UK); 0.2 mM dNTPs (Amersham-Pharmacia); 0.1 M GC-Melt (Clontech Laboratories, USA); and 0.5 units of *Taq* DNA polymerase (5 U/µl; Advanced Biotechnologies, UK).

The same PCR thermocycling conditions as described for the masterplate screening were utilized. The 96 individual PCR reactions were analysed on a 2% agarose gel containing ethidium bromide and a positive well was identified which gave the 102 bp PCR fragment.

**Colony PCR:** One ml of the glycerol stock from the positive sub-plate well was diluted 1:100 in Luria broth (LB). One ml and 10 ml of the aforementioned dilution were then plated on two separate agar plates containing Luria broth ("LB"), and 100 µg/ml carbenicillin. The LB plates were then incubated overnight at 30°C. From these plates, 96 colonies were picked into a new 96-well PCR plate containing: 0.2 mM of each of the two aforementioned gene-specific primers, 1x standard PCR buffer (Buffer V; Advanced Biotechnologies, UK), 0.2 mM dNTPs (Amersham-Pharmacia), 0.1 M GC-Melt (Clontech Laboratories, USA), and 0.5 units of *Taq* DNA polymerase (5 U/µl; Advanced Biotechnologies, UK) in a final volume of 25 µl.

The same PCR thermocycling conditions as described for the masterplate screening were utilized. The 96 individual PCR reactions were then analysed on a 2% agarose gel containing ethidium bromide. A positive colony containing the 102 bp fragment was subsequently identified.

Sequencing of the plasmid DNA prepared from this positive colony revealed a full-length cDNA of 861 bp (SEQ ID NO:8). The cDNA coded for a pre-pro-neublastin (SEQ ID NO:9). Automated DNA sequencing was performed using the BigDye® terminator cycle sequencing kit (PE Applied Biosystems, USA). The sequencing gels were run on the ABI Prism 377 (PE Applied Biosystems, USA).

#### **Method 2: Cloning Neublastin cDNA from Human Brain:**

An additional method of amplifying the full-length Neublastin cDNA or cDNA fragment can be performed by RACE (Rapid Amplification of cDNA ends) and the Neublastin-specific primers NBNint.sense and NBNint.antisense described above, combined with vector-

specific or adapter-specific primers, for example by using the Marathon cDNA amplification kit (Clontech Laboratories, USA, Cat. No. K1802-1).

Whole human brain Marathon-Ready cDNA (Clontech Laboratories, USA, Catalogue. No. 7400-1) can be used to amplify the full-length neublastin cDNA. Useful primers for  
 5 amplification include a neublastin top strand primer 5'-ATGGAAGTTGGACTTGG-3' (SEQ ID NO:19) ("NBNext.sense"), and a neublastin bottom strand primer 5'-TCCATCACCCACCGGC-3' (SEQ ID NO:20) ("NBNext.antisense"), combined with the adaptor primer AP1 included with the Marathon-Ready cDNA. An alternative top strand primer has also been used, 5'-CTAGGAGCCCATGCCC-3' (SEQ ID NO:28). A further  
 10 alternative bottom strand primer, 5'-GAGCGAGCCCTCAGCC-3' (SEQ ID NO:33) may also be used. Likewise, alternative bottom strand primers SEQ ID NOS.: 24 and 26 may also be used.

#### **Method 3: Cloning neublastin cDNA from Human Brain:**

Another method of cloning neublastin cDNA is by screening human adult or fetal brain  
 15 libraries with one or more neublastin probes described herein (and as exemplified in Figure 1). These libraries include:  $\lambda$ gt11 human brain (Clontech Laboratories, USA, Cat. No. HL3002b); or  $\lambda$ gt11 human fetal brain (Clontech Laboratories, USA, Cat. No. HL3002b).

#### **Method 4: Rapid-Screening of Mouse Fetal cDNA for the neublastin Gene**

20 A rapid screening procedure for the neublastin gene was performed in the following manner. A 96-well master plate, containing plasmid DNA from 500,000 cDNA clones, was loaded with approximately 5000 clones per well. A 96-well sub-plate was utilized with *E. Coli* glycerol stock containing 50 clones per well. Three rounds of PCR-mediated amplification was performed in order to identify a gene of interest (*i.e.*, neublastin).

25 **Master Plate Screening:** A mouse fetal cDNA master plate was screened by 96-well PCR using gene-specific primers to isolate the mouse neublastin cDNA. The following two primers were synthesised:

(1) neublastin C2 primer (NBNint.sense): 5'-GGCCACCGCTCCGACGAG-3' (SEQ ID NO:21); and (2) neublastin C2as primer (NBNint.antisense):  
 30 5'-GGCGGTCCACGGTTCTCCAG-3' (SEQ ID NO:22). By using these two gene-specific primers a 220 bp positive PCR product was identified. The 220 bp nucleic acid possessed the following sequence (SEQ ID NO:14):

5' -GGCCACCGCTCCGACGAGCTGATACGTTTCCGCTTCTGCAGCGGCTCGTG  
 CGCCGAGCACGCTCCCAGCACGATCTCAGTCTGGCCAGCCTACTGGGCGCTGG  
 GGCCCTACGGTCGCCTCCCGGGTCCCGGCCGATCAGCCAGCCCTGCTGCCGGC  
 CCACTCGCTATGAGGCCGTCTCCTTCATGGACGTGAACAGCACCTGGAGAACC  
 5 GTGGACCGCC-3'

96-well PCR reactions were then performed in the following manner. Thirty nanograms of mouse fetal brain cDNA (6 ng/μl; Origene Technologies) was obtained from the corresponding well of the master plate and placed in a total volume of 25 μl which also contained: 0.2 mM of each of the two aforementioned gene-specific primers (*i.e.*, C2 primer (NBNint.sense) and neublastin C2as primer (NBNint.antisense)), 1x standard PCR buffer (Buffer V; Advanced Biotechnologies, UK), 0.2 mM dNTPs (Amersham-Pharmacia), 0.1 M GC-Melt (Clontech Laboratories, USA), and 0.5 units of *Taq* DNA polymerase (5 U/μl; Advanced Biotechnologies, UK).

The following PCR thermocycling conditions were utilized: an initial denaturation at 94°C for 3 minutes, followed by 35 cycles of denaturation at 94°C for 1 minute each, annealing at 55°C for 1 minute, extension at 72°C for 90 seconds; and a final extension at 72°C for 5 minutes. The 96 individual PCR reactions were analysed on a 2% agarose gel containing ethidium bromide stain. The 220 bp, positive PCR product seen from a well was found to correspond to a unique 96-well sub-plate. The 96 individual PCR reactions were then analysed by gel electrophoresis on a 2% agarose gel containing ethidium bromide stain. The 220 bp positive PCR product which had been identified corresponded to a unique well of the 96-well sub-plate.

**Sub-Plate Screening:** The 96-well mouse fetal sub-plate was screened by PCR-mediated amplification by placing 1 μl of the glycerol stock from the corresponding sub-plate well into a final, total volume of 25 μl which contained: 0.2 mM of each of the two aforementioned gene-specific primers; 1x standard PCR buffer (Buffer V; Advanced Biotechnologies, UK); 0.2 mM dNTPs (Amersham-Pharmacia); 0.1 M GC-Melt (Clontech Laboratories, USA); and 0.5 units of *Taq* DNA polymerase (5 U/μl; Advanced Biotechnologies, UK). The PCR thermocycling was performed according to the conditions described above for the master plate screening.

The individual 96 PCR reactions were then analysed on a 2% agarose gel containing ethidium bromide and a positive well was identified which produced the 220 bp fragment.

**Colony PCR:** One ml of the glycerol stock from the positive sub-plate well was diluted 1:100 in Luria broth (LB). One ml and 10 ml of the aforementioned dilution were then plated on two separate LB plates, containing 100 µg/ml carbenicillin, and incubated at 30°C overnight. A total of 96 colonies were isolated and transferred to a 96-well PCR plate containing: 0.2 mM of each of the two aforementioned gene-specific primers, 1x standard PCR buffer (Buffer V; Advanced Biotechnologies, UK), 0.2 mM dNTPs (Amersham-Pharmacia); 0.1 M GC-Melt (Clontech Laboratories, USA), and 0.5 units of *Taq* DNA polymerase (5 U/µl; Advanced Biotechnologies UK) in a final volume of 25 µl.

PCR thermocycling was performed according to the conditions described above (*see*, "master plate screening", *infra*). The 96 individual PCR reactions were analysed by gel electrophoresis on a 2% agarose gel containing ethidium bromide. A positive colony was identified by the presence of the 220 bp fragment. Plasmid DNA was prepared from this positive colony. The clone was sequenced by automated DNA sequencing using the BigDye® terminator cycle sequencing kit with AmpliTaq DNA polymerase. The sequencing gels were run on the ABI Prism 377 (PE Applied Biosystems). The resulting sequence of this clone revealed a full-length cDNA of 2136 bp (SEQ ID NO:15). The cDNA includes an open reading frame with the predicted amino acid sequence shown in SEQ ID NO:16, which codes for a mouse pre-pro-neublastin polypeptide.

## 20 **Example 2: Cloning of Genomic Neublastin**

As discussed above, applicants identified a 290 bp nucleic acid fragment in two human BAC clones with entries in GenBank (with the Accession Nos. AC005038 and AC005051) which had regions of homology to persephin and to the flanking sequences of persephin. Applicants used the 861 bp predicted sequence described above to design additional primers, with the goal of cloning a nucleic acid encoding additional Neublastin nucleic acids using Lasergene Software (DNASTar, Inc.). Two pairs of primers were used to clone the Neublastin gene by using PCR reactions on genomic DNA. The two pairs of primers are illustrated below.

### **Primer Pair No. 1**

5' CCA AgC CCA CCT ggg TgC CCT CTT TCT CC 3' (sense) (SEQ ID NO:23).  
 30 5' CAT CAC CCA CCg gCA ggg gCC TCT CAg 3' (antisense) (SEQ ID NO:24).

### **Primer Pair No. 2**

5' gAgCCCAtgCCCggCCTgATCTCAgCCCgA ggACA 3' (sense) (SEQ ID NO:25).

5' CCCTggCTgAggCCgCTggCTAgTgggACTCTgC 3' (antisense) (SEQ ID NO:26).

Using primer pair No. 1, a 887 bp DNA fragment was amplified from a preparation of human genomic DNA purchased from Clontech Laboratories, (Cat. No. 6550-1).

**PCR protocol:** PCR was performed using the Expand<sup>TM</sup> High Fidelity PCR system (Boehringer Mannheim) with buffer 1. The PCR reaction mixture was supplemented with 5 % dimethylsulfoxide (DMSO) and 17.5 pmol of each dNTP in a total volume of 50 µl. Thermocycling was performed with a pre-denaturation step at 94°C for 2 minutes, followed by 35 two-step cycles at 94°C for 10 seconds, and 68°C for 1 minute, respectively. Thermocycling was terminated by incubation at 68°C for 5 minutes. Thermocycling was carried out in a PTC-225 DNA Engine Tetrad thermocycler (MJ Research, MA). The PCR products were analysed by gel electrophoresis on 2% agarose (FMC) and then photographed.

The 887 bp fragment amplified from human genomic DNA with primer pair No. 1 was cloned into the pCRII vector (Invitrogen), and transformed into XL 1-Blue competent *E. coli* cells (Stratagene). The resulting plasmid, designated neublastin-2, was sequenced using Thermosequenase (Amersham Pharmacia Biotech). Sequencing products were analysed by electrophoreses on an ALFExpress automated sequencer (Amersham Pharmacia Biotech).

Fragments obtained by PCR amplification of human genomic DNA with the second pair of primers (Primer Pair No. 1, above), were sequenced, revealing an additional 42 bp region at the 3' prime end of the open reading frame. The full-length sequence was analysed by comparing it to the sequences of nucleic acids of other neurotrophic factors, as well as by mapping exon-intron boundaries using gene-finding software programs which identify probable splice junctions and regions of high coding potential using Netgene and Gene Mark software (Brunak et al., *J. Mol. Biol.*, 220, pp. 49-65 (1991); Borodovsky et al., *Nucl. Acids Res.*, 23, pp. 3554-62 (1995)). The exon-intron boundaries were confirmed by the cDNA obtained from the Rapid Screen described above.

As illustrated in FIG. 7, the resulting neublastin gene has two exons separated by a 70 bp intron. Together, the exons have a predicted amino acid sequence of a full-length Neublastin polypeptide. The predicted cDNA (SEQ ID NO:3) contains an open reading frame (ORF) encoding 238 amino acid residues (SEQ ID NO:4). The Neublastin-2 clone contained the complete coding sequence of pro-neublastin. The amino acid sequence encoded by the gene showed high homology to three proteins, persephin, neurturin, and GDNF.



**Example 3: Expression of Neublastin Nucleic Acids**

Expression of Neublastin RNA was detected in both nervous and non-nervous tissue in rodents and in humans, and at various developmental immature and adult stages, using the techniques described below.

- 5           **Method of detecting Neublastin RNA expression using RT-PCR:** Based on the Neublastin DNA sequence identified as SEQ ID NO:1, the following primers were synthesised: (1) a Neublastin C2 primer 5'-GGCCACCGCTCCGACGAG-3' (SEQ ID NO:21), and (2) a Neublastin C2as primer 5'-GGCGGTCCACGGTTCTCCAG-3' (SEQ ID NO:22). This primer set was used to RT-PCR amplify a DNA fragment from adult and fetal human whole-brain mRNA. Among the DNA fragments produced by this reaction was one of 220 bp. Identification of this 220 bp DNA fragment confirmed that the Neublastin gene is expressed in adult and fetal brain tissue. A 220 bp DNA fragment was also amplified from genomic DNA with using these primers.

**Method of detecting Neublastin RNA expression by northern blot hybridization:**

- 15 Northern blots with polyA<sup>+</sup> RNA from adult human tissue were purchased from a commercial supplier (Clontech Laboratories, USA) and probed with a <sup>32</sup>P-labeled Neublastin cDNA. The labelled Neublastin cDNA was prepared according to the methods described in Example 1, above.

- 20           **Preparation of Probes:** A Neublastin nucleic acid DNA fragment (nucleotides 296-819 of SEQ ID NO:8) was labelled by the Rediprime II labelling kit (Amersham; Cat. No. RPN1633) for use as a hybridization probe, as recommended by the manufacturer. Briefly, the DNA sample was diluted to a concentration of 2.5-25 ng in 45 µl of 10 mM TE Buffer (10 mM Tris-HCl, pH 8.0, 1 mM EDTA). The DNA was then denatured by heating the sample to 95-100°C for 5 minutes in a boiling water bath, quick cooling the sample by placing it on ice for 5 minutes, and then briefly centrifuging it to bring the contents to the bottom of the reaction tube. The total amount of denatured DNA was added together with 5 µl of Redivue (<sup>32</sup>P) dCTP (Amersham Pharmacia Biotech Ltd.) in the reaction tube containing buffered solution of dATP, dGTP, dTTP, exonuclease free Klenow enzyme and random primer in dried stabilised form. The solution was mixed by pipetting up and down 2 times, moving the pipette tip around in the solution, and the reaction mixture was incubated at 37°C for 10 minutes. The labelling reaction was stopped by adding 5 µl of 0.2 M EDTA. For use as a hybridization probe the labelled DNA was denatured to single strands by heating the DNA sample to 95-100°C for 5

minutes, then snap cooling the DNA sample on ice for 5 minutes. The tube was centrifuged and its contents mixed well. Finally the single-stranded DNA probe was purified using the Nucleotide Removal Kit (Qiagen).

**Hybridization Techniques:** Prepared northern blots were purchased from a commercial supplier ("Multiple Tissue Northern Blots, Clontech Laboratories, USA, Catalogue Nos. 7760-1 and 7769-1) and were hybridized according to the manufacturer's instructions using the neublastin  $^{32}\text{P}$ -labeled probe prepared above. For hybridization, ExpressHyb Solution (Clontech Laboratories, USA) was used, and a concentration of approximately 3 ng/ml of the labelled probe was employed. The ExpressHyb solution was heated to 68°C and then stirred to dissolve any precipitate. Each northern blot membrane (10x10 cm) was pre-hybridized in at least 5 ml of ExpressHyb Solution at 68°C for 30 minutes in a Hybaid Hybridization Oven according to the manufacturer's instructions. The neublastin  $^{32}\text{P}$ -labeled probe was denatured at 95-100°C for 2 minutes and then chilled quickly on ice. Fourteen microliters (14  $\mu\text{l}$ ) of the labelled probe was added to 5 ml of fresh ExpressHyb, and thoroughly mixed. The ExpressHyb Solution used in the pre-hybridization was replaced by evenly distributing over the blots the 5 ml of fresh ExpressHyb Solution containing labelled DNA probe. Blots were incubated at 68°C for 1 hour in a Hybaid hybridization Oven. After incubation, the blots were rinsed and washed several times at low stringency (2x SSC buffer containing 0.05 % SDS at room temperature) followed by a high stringency wash (0.1x SSC containing 0.1 % SDS at 50°C) (20X SSC is 0.3 M NaCl/0.3 M Na citrate, pH 7.0). The blots were exposed to a Hyperfilm MP (Amersham Pharmacia Biotech Ltd.) at -80°C using intensifying screens.

The results of the northern blot hybridization experiments are presented in FIG. 1. FIG. 1A (left) and FIG. 1B (right) are northern blots of polyA<sup>+</sup> RNA which were probed with  $^{32}\text{P}$ -labelled neublastin cDNA as described in Example 3. The markers represent polynucleotides of 1.35 kilobase pairs ("kb"), 2.4 kb, 4.4 kb, 7.5 kb, and 9.5 kb in size. The membrane of FIG. 1A was prepared with mRNA extracted from various adult human tissues: From the results of the northern blot hybridization analysis, applicants conclude that neublastin mRNA is expressed in many adult human tissues. The highest level of neublastin expression is detected in the heart, in skeletal muscle and in the pancreas. The membrane of FIG. 1B was prepared with RNA extracted from various regions of the adult human brain. Within the adult brain, the highest level of expression is seen in the caudate nucleus and in the thalamus. An mRNA

transcript of approximately 5 kb was the predominant form of neublastin mRNA expressed in the brain.

**Method of detecting Neublastin RNA expression using by in situ Hybridization in Tissues:**

The following techniques are used to measure the expression of neublastin RNA in animal tissues, *e.g.*, rodent tissues, with a neublastin anti-sense probe.

**Expression in mice:**

**Preparation of Tissue Samples:** Time pregnant mice (B&K Universal, Stockholm, Sweden) were killed by cervical dislocation on gestational day 13.5 or 18.5. Embryos were removed by dissection under sterile conditions, and immediately immersed in a solution of 0.1M phosphate buffer (PB) containing 4% paraformaldehyde ("PFA") for 24-30 hours, and then removed from the PFA and stored in PBS. The tissue was prepared for sectioning by immersing the tissue in a solution of 30% sucrose, and then embedding it in TissueTech (O.C.T. Compound, Sakura Finetek USA, Torrance, CA). Six series of coronal or sagittal sections (12  $\mu$ m each) were cut on a cryostat and thaw mounted onto positively charged glass slides. Neonatal heads/brains (P1, P7) were fixed following the same protocol as for the embryonic stages, and adult brain tissue was dissected, immediately frozen on dry ice, and cut on a cryostat without any prior embedding.

**Preparation of Neublastin Riboprobes:** An antisense neublastin RNA probe (hereafter a "neublastin riboprobe") was made as follows. Nucleotides 1109-1863 of the mouse neublastin cDNA sequence (SEQ ID NO:15) were sub-cloned into the BlueScript vector (Stratagene). The resulting plasmid was cut into a linear DNA using *EcoRI* restriction endonuclease. The *EcoRI* DNA fragment was *in vitro* transcribed with T3 RNA polymerase and the digoxigenin ("DIG") RNA Labelling Kit according to the manufacturer's instructions (Boehringer Mannheim).

**Hybridization:** Cryostat sections were fixed for 10 minutes in 4% PFA, treated for 5 minutes with 10 mg/ml of proteinase K, dehydrated sequentially in 70% and 95% ethanol for 5 and 2 min, respectively, and then allowed to air dry. Hybridization buffer (50% deionized formamide, 10% of a 50% dextran sulphate solution, 1% Denhardt's solution, 250  $\mu$ g/ml yeast tRNA, 0.3M NaCl, 20mM Tris-HCl (pH8), 5mM EDTA, 10mM NaPO<sub>4</sub>, 1% sarcosyl) containing 1  $\mu$ g/ml of the DIG-labelled probe was heated to 80°C for 2 minutes and applied onto the sections. The sections was then covered with parafilm and incubated at 55°C for 16-18 hours.

The next day the sections were washed at high stringency (2x SSC containing 50% formamide) at 55°C for 30 minutes, and then washed in RNase buffer and incubated with 20µg/ml of RNaseA for 30 minutes at 37°C. In order to detect the DIG-labelled probe, sections were pre-incubated in blocking solution (PBS containing 0.1% Tween-20 and 10% heat-inactivated goat serum) for 1 hour and then incubated over night at 4°C with a 1:5000 dilution of alkaline-phosphatase-coupled anti-DIG antibody (Boehringer Mannheim). The following day, each section was given four, two-hour washes in PBS containing 0.1% Tween-20, and then given two ten-minute washes in NTMT buffer (100 mM NaCl, 100 mM Tris-HCl (pH9.5), 50 mM MgCl<sub>2</sub>, 0.1% Tween-20). The sections were then incubated in BM-purple substrate containing 0.5mg/ml of levamisole for 48 hours. The color reaction was stopped by washing in PBS. The sections were air dried and covered with cover-slip with DPX (KEBO-lab, Sweden).

The results of the in situ hybridization reactions are presented in Table 1.

Table 1: Expression of neublastin in Mice

Structure	E13.5	E18.5	P1	P7	Adult
Forebrain	++				
Ventral Midbrain	-				
Dorsal Root ganglia	++				
Spinal chord	+				
Retina		+++	+++	+	
Olfactory bulb		++	++	++	
Tooth pulp		++	++	+	
Trigeminal ganglia		++	++	++	
Striatum		+	+	++	
Cortex		++	++	++	+
Dentate gyrus				++	+

As shown in Table 1, at embryonic day 13.5 ("E13.5"), neublastin was expressed in the spinal chord and in the hindbrain, and weakly in the forebrain. Neublastin expression was also detected in the developing retina and in the sensory ganglia (dorsal root ganglia and trigeminal ganglia (V)). Outside the nervous system, a weak signal was found in the kidney, the lung and the intestine, indicating that neublastin is also expressed in those tissues.

At embryonic day 18.5 ("E18.5"), neublastin was expressed most prominently in the trigeminal ganglion (V). Neublastin expression was also detected in the retina, the striatum, and the cortex. In addition, expression was seen in tooth anlage.

Again referring to Table 1, increased neublastin expression, from the E18.5 time-point to postnatal days 1 and 7, was seen in the cortex, the striatum and the trigeminal ganglion (V). Neublastin expression was more prominent in the outer layers of the cortex than in the inner layers of the cortex. On P7, expression was found in the same structures as at day 1 but in addition neublastin expression was found in the hippocampus, especially in the dentate gyrus and in the cerebellum. In the adult murine brain, neublastin was strongly expressed in dentate gyrus, with very low or undetectable levels of neublastin expression detected other tissues tested.

#### **Expression in Rat:**

The following experiment describes the hybridization of rat tissues with a alkaline-phosphatase- labelled oligodeoxynucleotide neublastin anti-sense probe.

**Preparation of tissue samples:** Rat embryos (E14) were obtained from pregnant Wistar rats (Møllegaard, Denmark) following pentobarbital anaesthesia. Postnatal rats (P0, P7, adult) were killed by decapitation. Dissected brains and whole heads were immediately immersed in cold 0.9% NaCl, fresh frozen and sectioned at 20 µm on a cryostat (coronal and sagittal sections, 10 series).

**In situ hybridization:** Two series of sections were hybridized using an anti-sense alkaline-phosphatase (AP) conjugated oligodeoxynucleotide probe (5'-NCA GGT GGT CCG TGG GGG GCG CCA AGA CCG G-3' (SEQ ID NO:27), Oligo. No. 164675, DNA Technology, Denmark.). This probe is complementary to bases 1140 to 1169 of the mouse neublastin cDNA of SEQ ID NO:15).

Prior to hybridization, the sections were air dried at room temperature, heated at 55°C for 10 min., and then treated with 96% ethanol at 4°C overnight. The sections were then air dried and incubated in hybridization medium (5.0 pmol probe/ml) overnight at 39 °C (*Finsen et al.*, 1992, *Neurosci.* 47:105-113; *West et al.*, 1996, *J. Comp. Neurol.*, 370:11-22).

Post-hybridization treatment consisted of four, thirty-minute rinses in 1x SSC (0.15M NaCl, 0.015 M Na-citrate) at 55 °C, followed by three ten-minute rinses in Tris-HCl, pH 9.5 at room temperature prior to applying AP developer. AP developer was prepared immediately before use and contained nitroblue tetrazoleum (NBT, Sigma), 5-bromo, 4-chloro, 3-indolylphosphate (BCIP, Sigma), and Tris-HCl-MgCl<sub>2</sub> buffer, pH 9.5 (*Finsen et al.*, *Neurosci.* 1992 47 105-113). AP development took place in the dark at room temperature for 48 hours. The color reaction was stopped by rinsing the sections in distilled water. The sections were

dehydrated in graded acetone, softened in xylene-phenol creosote (Allchem, UK), cleared in xylene, and coverslipped using Eukitt (Bie & Berntsen, Denmark).

Control reactions consisted of (1) pre-treating the sections with RNase A (50 µg/ml, Pharmacia, Sweden) prior to hybridization; (2) hybridizing the sections with a hundred-fold excess of unlabelled probe; and (3) hybridizing the sections with hybridization buffer alone. The results of the hybridization reactions are presented in Table 2.

**Table 2: Expression of neublastin in rats**

Structure	E14	P0/P1	P7	Adult
Forebrain	++			
Ventral Midbrain	-			
Dorsal root ganglia	++			
Spinal cord	+			
Retina	+			
Olfactory bulb	(+)	++	++	
Cerebellum		+	++	+
Trigeminal ganglia		++	++	
Striatum		+	+(+)	
Cortex	(+)	++	++	+
Hippocampus		(+)	++	++

At embryonic day 14 (E14), neublastin was weakly expressed in rat embryos in the forebrain, in the hindbrain, and in the spinal cord. Neublastin mRNA was also detected in the eye (retina), dorsal root ganglia, the trigeminal ganglia (V), and in the kidneys, lungs, heart, liver, and intestines. In newborn (P0) rats there was marked neublastin expression in the cortex and in the striatum. Neublastin expression was also detected in the olfactory bulb and in the hippocampus. In 7-day-old (P7) rats, neublastin was expressed in the cortex, the striatum, the olfactory bulb, and in the cerebellum. A marked signal was seen in the hippocampus. In adult rats, very low or undetectable levels of neublastin expression were detected in most areas of the brain. Weak signals were detected in the thalamic nucleus, and marked neublastin expression was detected in the hippocampus.

#### **Example 4: Neublastin Polypeptides**

The open reading frame, or coding region (CDS), identified in SEQ ID NO:8 encodes the pre-pro-polypeptide (designated "pre-pro-neublastin"). The amino acid sequence predicted from this open reading frame is shown in SEQ ID NO:9. Based on SEQ ID NO:9, three variants of neublastin polypeptides were identified. These variants include: (i) the polypeptide

designated herein as NBN140, which possesses the amino acid sequence designated as SEQ ID NO:10; (ii) the polypeptide designated herein as NBN116, which possesses the amino acid sequence designated as SEQ ID NO:11; and (iii) the polypeptide designated herein as NBN113, which possesses the amino acid sequence designated as SEQ ID NO:12.

- 5           Similarly, based on the coding region (CDS) as identified in SEQ ID NO:3, which encodes the pre-pro-polypeptide possessing the amino acid sequence (designated as SEQ ID NO:4), three variants of neublastin were identified. These variants include: (i) the polypeptide which possesses the amino acid sequence designated as SEQ ID NO:5; (ii) the polypeptide which possesses the amino acid sequence designated as SEQ ID NO:6; and (iii) the polypeptide
- 10   which possesses the amino acid sequence designated as SEQ ID NO:7.

Based on a Clustal W (1.75)-based multiple sequence alignment, neublastin of SEQ ID NO:9 (line 2) was aligned with the amino acid sequences of neurturin (SEQ ID NO:49; line 1), persephin (SEQ ID NO:50; line 3) and GDNF (SEQ ID NO:51; line 4). This alignment is illustrated in Table 3.

Table 3:

## Amino Acid Sequence Comparison of Neublastin to Persephin, Neurturin, and GDNF

Neurturin-full	-----MQRWKAALASVLCSSVLSIWMCREGLLLSHRLGPA
Neublastin	MELGLGGLSTLSHCPWPRRQPALWPTLAALALLSSVAEASLGSAPRSPAPREGPPP
Persephin-full	-----
GDNF_HUMAN-full	----MKLWDVVAVCLVLLHTASAFPLPAGKRPPEAPAEDRSLGRRRAPFALSSDS
Neurturin-full	LVPLHRLPRTLDARIARLAQYRALLQGAPDAMELRELTWPAGRPPGPRRRAGPRRR
Neublastin	VLASPAHLPGGRTARWCSSGRARRPPPPQPSRPAPPPPPAPPALPRGGRAARAGGPG
Persephin-full	-MAVGKFLLGSLLLSLQLGQGWGPDARGVPVADGEFSSEQVAKAGGTWLGTNRPL
GDNF_HUMAN-full	NMPEDYPDQFDDVMDFIQATIKRLKRSPDKQMAVLPRRERNRQAAAANPENSRGKG
Neurturin-full	RARARLGARPCGLRELEVRVSELGLGYASDETVLFRYCAGACEA-AARVYDLGLRR
Neublastin	SRARAAGARGCRLRSQVFPVRAELGLGHRSEDLVRFRCGSGSCRR-ARSPHDLSLAS
Persephin-full	ARLRRALSGPCQLWSLTLSVAELGLGYASEEKVIFRYCAGSCPRGARTQHGLALAR
GDNF_HUMAN-full	RRGQRGKNRGCVLTAIHLNVTDLGLGYETKEELIFRYCAGSCDA-AETTYDKILKN
	* * : * ***** : * : * : * : * * * * *
Neurturin-full	LRQRRRLRRE---RVRAQPCCRPTAYEDEVSFLDAHSRYHTVHELARECACV-
Neublastin	LLGAGALRPPPGSRPVSQPCCRPTRYE-AVSFMDVNSTWRTVDRLSATACGCLG
Persephin-full	LQGQGRAHGG-----PCCRPTRYT-DVAFLDDRHRWQRLPQLSAAACGCGG
GDNF_HUMAN-full	LSRNRLVSD---KVGQACCRPIAFDDDLFLDDNLVYHILRKHSKRCGCI-
	* .***** : : : * . : : : * * *

\* indicates positions which have a single, fully conserved residue.

: indicates that one of the following 'strong' groups is fully conserved:

-STA, NEQK, NHQK, NDEQ, QHRK, MILV, MILF, HY, FYW.

. indicates that one of the following 'weaker' groups is fully conserved:

-CSA, ATV, SAG, STNK, STPA, SGND, SNDEQK, NDEQHK, NEQHRK, HFY.

From the amino acid sequence alignment shown in Table 3, it can be seen that neublastin has seven conserved cysteine residues at locations that are conserved within the TGF- $\beta$  superfamily. In one embodiment, the preferred neuroblastin polypeptide contains (seven) cysteines conserved as in SEQ ID NO:2 at positions 8, 35, 39, 72, 73, 101 and 103, or as in SEQ ID NOS: 4 and 9 at positions 43, 70, 74, 107, 108, 136 and 138. These seven conserved cysteine residues are known within the TGF-b superfamily to form three intramonomeric disulfide bonds (contemplated, e.g., in SEQ ID NO:2 between cysteine



residues 8-73, 35-101, and 39-103, and, *e.g.*, in SEQ ID NOS: 4 and 9 between cysteine residues 43-108, 70-136, and 74-138) and one intermonomeric disulfide bond (contemplated, *e.g.*, in SEQ ID NO:2 between cysteine residues 72-72, and, *e.g.*, in SEQ ID NOS: 4 and 9 between cysteine residues 107-107), which together with the extended beta strand region constitutes the conserved structural motif for the TGF- $\beta$  superfamily. See, *e.g.*, *Daopin et al.*, Proteins 1993 17 176-192.

Based on this sequence alignment, neublastin was shown to be a member of the GDNF subfamily of neurotrophic factors (LGLG - FR(Y/F)CSGSC - QxCCRP - SAxxCGC, the GDNF subfamily fingerprint, underlined in Table 3).

The truncated neublastin polypeptides described herein preferably include a polypeptide sequence that encompasses the seven cysteine residues conserved in the neublastin sequence. For example, the truncated neublastin polypeptides preferably include amino acids amino acids 15-113 (a 99 AA NBN form) of a NBN113 polypeptide, or amino acids 12-113 of the NBN113 polypeptide (a 102 AA NBN form). These amino acid sequences can be found at, *e.g.*, amino acids 42-140 of the human NBN polypeptide sequence shown in SEQ ID NO:9 (99 AA NBN polypeptide; SEQ ID NO:48); and amino acids 39-140 of SEQ ID NO:9 (102 AA NBN polypeptide; SEQ ID NO:45), respectively. The sequences are also found at, *e.g.*, amino acids 126-224 of SEQ ID NO:34 (rat 99 AA NBN polypeptide) and at amino acids 123-224 of SEQ ID NO:34 (rat 102 AA NBN polypeptide). Likewise, also included are the truncated neublastin sequences of NBN99, NBN100, NBN101, NBN102, NBN103, NBN104, NBN105, NBN106, NBN107, NBN108, NBN109, NBN110, NBN111 and NBN112, as defined above.

The homology of neublastin to other members of the GDNF family was calculated, and the results are presented Table 4, below.

**Table 4: Homology of Neublastin Polypeptides to other members of the GDNF Family**

	Mature Protein NBN140			Mature Protein NBN113			
	Homology		Homology of full length peptides	Homology		Homology of full length peptides	
Neurotrophic Factor	Identity	Overlap (aa)	Strong Homology	Identity	Identity	Overlap (aa)	Strong Homology
GDNF	34%	137	48%	31.9%	36%	111	52%
	(47/137)		(67/137)		(41/111)		(59/111)

NTN	48%	127	56%	36.9%	49%	114	57%	44.7%
	(61/127)		(72/127)		(56/114)		(66/114)	
PSP	44%	125	56%	36.9	45%	111	57%	44.3%
	(55/125)		(71/125)		(51/111)		(65/111)	
IHA	31%	81	-	25.2%	31%	81	-	22.5%
	(25/81)				(25/81)			
TGF- $\beta$ 2	23%	73	-	18.5%	23%	73	-	20.2%
	(17/73)				(17/73)			

GDNF = Glial cell line Derived Neurotrophic Factor

NTN = Neurturin

PSP = Persephin

5 IHA = Inhibin- $\alpha$

TGF- $\beta$ 2 = Transforming Growth Factor- $\beta$ 2

Strong homology indicates that one of the following "strong" groups are conserved: STA, NEQK, NHQK, NDEQ, QHRK, MILV, MILF, HY, FYW.

#### 10 **Example 5: Production of Neublastin**

Neublastin has been produced in both eukaryotic and prokaryotic cells, as described below.

**Expression Vectors** The cDNA encoding neublastin was inserted into the eukaryotic expression vector pUbi1Z. This vector was generated by cloning the human UbC promoter into  
 15 a modified version of pcDNA3.1/Zeo. The unmodified pcDNA3.1/Zeo is commercially available (Invitrogen). The modified pcDNA3.1/Zeo is smaller than the parent vector, because the ampicillin gene (from position 3933 to 5015) and a sequence from position 2838 to 3134 were removed. In this modified version of pcDNA3.1/Zeo, the CMV promoter was replaced with the UbC promoter from pTEJ-8 (*Johansen et al.*, FEBS Lett. 1990 267 289-294), resulting  
 20 in pUbi1Z.

**Mammalian Cell Expression** The pUbi1Z vector which contained neublastin coding sequences was then transfected into the mammalian cell line HiB5, which is an immortalised rat neural cell line (Renfranz et al., *Cell*, 66, pp. 713-729 (1991)). Several HiB5 cell lines stably expressing neublastin (as determined by RT-PCR) have been established. In one of  
 25 these stable cell lines, HiB5pUbi1zNBN22 expression was confirmed by hybridizing total RNA on a Northern blot with a <sup>32</sup>P-labelled neublastin probe. The results of these studies are shown in FIG. 2. HiB5pUbi1zNBN22 was then used as a source of neublastin for some studies of neublastin neurotrophic activity.

FIG. 2 shows the expression of neublastin cDNA in the HiB5pUbi1zNBN22 clone (*i.e.*,  
 30 Northern blot probed with <sup>32</sup>P-labelled neublastin cDNA of the present invention as described *infra*). The blot was prepared by total RNA extracted from untransfected HiB5 cells,

HiB5pUbi1zNBN22 cells and HiB5pUbi1zGDNF14, respectively, as indicated. The positions of the 28S and 18S rRNA bands corresponding to 4.1 kb and 1.9 kb, respectively, are indicated on the blot.

As shown in FIG. 3, antibodies raised against neublastin-derived polypeptides also  
5 recognised a protein of approximately 13 kilodaltons ("kD") in conditioned medium from the HiB5pUbi1zNBN22 clone but not from non-transfected HiB5 cells (cf. Example 6).

The predicted molecular weights of the non-modified (*i.e.* lacking post-translational modifications) neublastin polypeptides NBN140 (SEQ ID NO:10), NBN116 (SEQ ID NO:11) and NBN113 (SEQ ID NO:12) were determined to be 14.7 kilodaltons ("kD"), 12.4 kD, and  
10 12.1 kD, respectively.

**Methods:** A Northern blot with total RNA (10 µg) from untransfected HiB5 cells and the HiB5pUbi1zNBN22 clone was prepared by electrophoresis on a 0.8 % formaldehyde agarose gel and blotted onto a nylon membrane (Duralone, Stratagene). The blot was hybridized and washed as described in Example 3 with a 1.3 kb <sup>32</sup>P-labelled probe prepared by  
15 random labelling covering SEQ ID NO: 8 and additional nucleotides from the 5'UTR and 3'UTR of the neublastin cDNA. The blot was exposed to a Hyperfilm MP (Amersham) at -80°C using intensifying screens.

Conditioned medium from Hib5pUbi1zNBN22, or untransfected Hib5 cells incubated overnight in serum-free medium supplemented with N2 supplement (Life Technologies; Cat.  
20 No. 17502-048) was concentrated and separated on 15% polyacrylamide gels (Amersham Pharmacia Biotech; Cat. No. 80-1262-01). Proteins were transferred to PVDF-membranes (Amersham Pharmacia Biotech; Cat. No. RPN-303F) and non-specific protein-binding sites were blocked with 5% non-fat dry milk in PBS with 0.1% Tween-20. Membranes were incubated overnight with a polyclonal neublastin antibody (1:1000), followed by incubation  
25 with a secondary anti-rabbit IgG antibody (Amersham Pharmacia Biotech; Cat. No. NA 934) conjugated to horseradish peroxidase (1:2000). Immunostaining was visualised using enhanced chemiluminescence (ECL) (Amersham Pharmacia Biotech; Cat. No. RPN2109) or ECL+ (Amersham Pharmacia Biotech; Cat. No. RPN2132) according to the manufacturer's instructions (Amersham).

30 The results of these experiments are shown in Figure 3. Figure 3A and 3B are illustrations of the expression of neublastin protein in transfected HiB5 cells. Overnight medium from non-transfected HiB5 cells (Lane 1), or from an HiB5 clone stable transfected

with neublastin cDNA (Lane 2), were concentrated as described *infra*. The medium was then analyzed by Western blotting using two different polyclonal antibodies, Ab-1 and Ab-2 described in Example 10, specific for neublastin. In the medium derived from transfected cells, both of the antibodies were found to recognize a protein with a molecular weight of approximately 15 kDa. This protein was not seen in non-transfected HiB5 cells.

The cloned cDNA encoding neublastin can also be inserted into other eukaryotic expression vector, *e.g.*, the eukaryotic expression vector TEJ-8 (*Johansen et al.*, FEBS Lett., 1990, 267:289-294) or pcDNA-3 (Invitrogen), and the resulting expression plasmid transfected into an alternative mammalian cell line, *e.g.*, Chinese Hamster Ovary ("CHO") cells, the HEK293, the COS, the PC12, or the RN33b cell lines, or a human neural stem cell. Stable cell lines expressing neublastin are used, *e.g.*, to produce the neublastin protein.

### Expression in CHO Cells

**Construction of plasmid pJC070.14** In order to express the neublastin cDNA in Chinese hamster ovary cells, a cDNA fragment encoding the prepro form of human neublastin was inserted into the mammalian expression vector pEAG347 to generate plasmid pJC070.14. pEAG347 contains tandem SV40 early and adenovirus major late promoters (derived from plasmid pAD2beta; Norton and Coffin, *Mol. Cell. Biol.* 5: 281 (1985)), a unique NotI cloning site, followed by SV40 late transcription termination and polyA signals (derived from plasmid pCMVbeta; MacGregor and Caskey, *Nucl. Acids. Res.* 17: 2365 (1989)). In addition, pEAG347 contains a pUC19-derived plasmid backbone and a pSV2dhfr-derived dhfr for MTX selection and amplification in transfected CHO cells.

Plasmid pJC070.14 was generated in two steps. First, a fragment encoding the prepro form of human neublastin was isolated from plasmid pUbi1Z-NBN using the polymerase chain reaction with oligonucleotides KD2-824 5'AAGGAAAAAA GCGGCCGCCA TGGAACCTGG ACTTGGAGG3' (SEQ ID NO:31), KD2-825 5'TTTTTTCCTT GCGGCCGCT CAGCCCAGGC AGCCGCAGG3' (SEQ ID NO:32) and PFU polymerase. The fragment was cloned into the Srf-1 site of pPCR-Script Amp SK(+) to generate the plasmid pJC069. In the second step, a partial Not-1 digest was performed on plasmid pJC069 to generate a 685bp fragment (containing the neublastin gene) which was cloned into the Not-1 site of plasmid pEAG347 to generate plasmid pJC070.14. Transcription of the neublastin gene in plasmid pJC070.14 is controlled by the adenovirus major late promoter.

**Generation of CHO cell lines expressing Neublastin.** 200 µg of pJC070.14 was linearized by digestion with the restriction endonuclease Mlu-1. The DNA was extracted with phenol: chloroform: isoamyl alcohol (25:24:1) and ethanol precipitated. The linearized DNA was resuspended in 20mM Hepes pH7.05, 137mM NaCl, 5mM KCl, 0.7mM Na<sub>2</sub>HPO<sub>4</sub>, 6mM dextrose (HEBS) and introduced into ~4E7 CHO dukx B1(*dhfr*<sup>-</sup>) cells (p23) by electroporation (280V and 960 µF). Following electroporation, the cells were returned to culture in α+ Modified Eagle's Medium (MEM) supplemented with 10% fetal bovine serum (FBS) for two days. The cells were then trypsinized and replated in 100mm dishes (100,000 cells/plate) in α-MEM (lacking ribo- and deoxyribonucleosides), supplemented with 10% dialyzed FBS, for five days. The cells were subsequently split at a density of 100,000 cells/100mm plate, and selected in 200nM methotrexate. Resistant colonies were picked and scaled up to 6 well plates; conditioned media from each clone was screened using a specific assay for neublastin described below. The twelve clones expressing the highest level of neublastin were scaled up to T162 flasks and subsequently reassayed. As shown in Figure 10, the CHO cell lines produced neublastin in the range of 25 to 50 ng/ml.

**Ternary complex assay for neublastin.** The presence of neublastin was assessed in the media of CHO cell line supernatants using a modified form of a ternary complex assay described by Sanicola et al. (Proc Natl Acad Sci USA 94: 6238 (1997)).

In this assay, the ability of GDNF-like molecules can be evaluated for their ability to mediate binding between the extracellular domain of RET and the various co-receptors, GFRα1, GFRα2, and GFRα3. Soluble forms of RET and the co-receptors were generated as fusion proteins. A fusion protein between the extracellular domain of rat RET and placental alkaline phosphatase (RET-AP) and a fusion protein between the extracellular domain of rat GFRα1 (disclosed in published application WO9744356; November 27, 1997, herein incorporated by reference) and the Fc domain of human IgG1 (rGFRα1-Ig) have been described (Sanicola et al. Proc Natl Acad Sci USA 94: 6238 (1997)).

To generate a fusion protein between the extracellular domain of murine GFRα3 and the Fc domain of human IgG1 (mGFRα3-Ig), a DNA fragment encoding amino acids 1-359 of murine RETL3 was ligated to a fragment containing the Fc domain of human IgG1 and cloned into the expression vector pEAG347 to generate plasmid pGJ144. Plasmid pGJ144 was transfected into Chinese hamster ovary cells (CHO) to generate a stable cell line producing the fusion protein, which was purified on a Protein A Sepharose immunoaffinity column using

standard methods. In summary, if the GDNF-like molecule can mediate binding of the co-receptor to RET in this assay, then the RET-AP fusion protein will be retained on the plate and the amount that is retained can be measured using a chemiluminescent substrate for alkaline phosphatase.

- 5 Dynex Microlite-1 ELISA plates (Dynex Technologies) were coated with 1 µg/ml goat antibody specific for human Fc in 50mM bicarbonate/carbonate, pH 9.6 for 16hr. The plates were emptied and filled with 300 µl of 1% I-block (Tropix) in TBS/0.5% Tween-20 (TBST), for 1hr. After washing three times with TBST the wells were filled with 100 µl of 1 µg/ml rGFRα1-Ig or mGFRα3-Ig diluted in conditioned media from 293 EBNA cells expressing the  
10 RET-AP fusion gene. 100ul of conditioned media from the CHO neublastin clones was then added to the top well of a column of wells, and 2 fold serial dilutions were performed down each row of wells, and incubated for 1.5hr at room temperature. The plates were then washed three times with TBST, and twice with 200mM Tris pH9.8, 10mM MgCl<sub>2</sub> (CSPD buffer). The wash solution was then replaced with 425µM CSPD (Tropix) in CSPD buffer containing  
15 1mg/ml Sapphire chemiluminescence enhancer (Tropix), and incubated for 30' at room temperature. The chemiluminescent output was measured using a Dynatech luminometer.

- The initial experiments investigated whether neublastin produced by the CHO cell lines could mediate the binding of GFRα1 or GFRα3 to the extracellular domain of RET. As shown in Figure 11, conditioned medium from CHO cell clone #53 produced a robust signal in the  
20 ternary complex assay when the mGFRα3-Ig fusion protein was included, but no signal when the rGFRα1-Ig fusion protein was included, indicating that neublastin binds to GFRα3 but not to GFRα1. This behavior clearly distinguishes neublastin from GDNF; as shown in Figure 11, GDNF binds to GFRα1 but not to GFRα3. No signal was observed with either co-receptor fusion protein, when conditioned medium from the parental CHO cell line or straight medium  
25 was assayed.

- In order to quantify the expression levels of neublastin in the CHO cell lines, a standard curve was prepared using rGFRα1-Ig and GDNF starting at a concentration of 1ng/ml. Neublastin concentrations for the different CHO cell lines were then calculated using this standard curve; the levels produced by five CHO cell lines are shown in Figure 10. Because  
30 this estimation depends on the untested assumption that the binding affinity between GDNF and GFRα1 is similar to the binding affinity between neublastin and GFRα3, these levels are only approximate.

**Analysis of neublastin from CHO cell line supernatants.** In order to further analyze the neublastin produced by the CHO cell lines, the protein was extracted from the medium using the GFR $\alpha$ 3-Ig fusion protein and analyzed by western blots with polyclonal antibodies raised against neublastin peptides.

In the first experiment, the neublastin was extracted with mGFR $\alpha$ 3-Ig attached to Sepharose beads. mGFR $\alpha$ 3-Ig was attached to Sepharose beads using the conditions suggested by the manufacturer, Pharmacia Inc. 100  $\mu$ L of mGFR $\alpha$ 3-Ig-Sepharose was added to 1.0 mL samples of conditioned medium from a negative control CHO cell line or from the neublastin producing CHO cell line #16. The suspensions were incubated for two hours on a rocking platform. Each suspension was centrifuged and the supernatant removed followed with three 1.0 mL washes with 10 mM HEPES, 100 mM NaCl, pH 7.5. Each resin was resuspended in 100  $\mu$ L of 2X reducing sample buffer and heated to 100°C for 5 minutes. 20  $\mu$ L of the sample buffer supernatant and 10  $\mu$ L of a molecular weight standard (FMC) were applied to each well of a 10-20% precast SDS-PAGE gel (Owl Scientific). The gel was electrophoresed at 40 mA constant current for 72 minutes.

For western blot analysis, the protein was electroblotted to nitrocellulose (Schleicher and Schuell) in a Hofer Scientific apparatus in 10 mM CAPS, 10% methanol, 0.05% SDS, pH 11.2 buffer system (45 minutes at 400 mA constant current). After the transfer, the nitrocellulose filter was removed from the cassette and the molecular weight markers were visualized by staining with a solution of 0.1% Ponceau S in 1% acetic acid for 60 seconds. The membrane was cut into two sections and the excess stain was removed by gentle agitation in distilled water. The membranes were blocked in 2% nonfat dry milk in TBS overnight at 4°C. The membranes were incubated individually with two of the affinity-purified anti-neublastin peptide antibodies (R30 and R31) at a concentration of 1.0  $\mu$ g/mL in 2% nonfat dry milk in TBS. The membranes were washed with three 10 minute washes in TBS-Tween and incubated in a 1:5000 dilution of goat anti-rabbit IgG-HRP conjugate (Biorad) for 30 minutes. The membranes were washed with three 10 minute washes of TBS-Tween and developed with ECL substrate (Amersham). As shown in Figure 12, specific bands were detected in the proteins extracted from the neublastin producing CHO cell line with both antibodies (lanes 2 and 4), when compared to the bands observed in the extracted proteins from the negative control cell line (lanes 1 and 3).

The molecular weight of the lower species is about 13kD and probably represents the mature domain of neublastin, generated after cleavage of the pro- domain. This cleavage could occur after the fourth position Arg residues in any one of the three cleavage motifs (*e.g.*, -RXXR↓-) present in the prepro neublastin protein, to generate either the 140 AA, 116 AA or 113 AA forms, as set forth in SEQ ID NOS:10, 11, or 12, respectively. The predicted molecular weights of the non-modified (*i.e.*, lacking post-translational modifications) neublastin polypeptides NBN140 (SEQ ID NO:10), NBN116 (SEQ ID NO:11), and NBN113 (SEQ ID NO:12) were determined to be 14.7 kD, 12.4 kD, and 12.1 kD, respectively. Further analysis will be needed to confirm the structure of this species as well as the other neublastin specific bands.

In the second experiment, the neublastin was extracted with hGFR $\alpha$ 3-Ig captured on an ELISA plate. To generate a fusion protein between the extracellular domain of human GFR $\alpha$ 3 (disclosed in published application WO97/44356; November 27, 1997, herein incorporated by reference) and the Fc domain of human IgG1 (hGFR $\alpha$ 3-Ig), a DNA fragment encoding amino acids 1-364 of human GFR $\alpha$ 3 was ligated to a fragment containing the Fc domain of human IgG1 and cloned into the expression vector CH269 described by Sanicola et al. (Proc Natl Acad Sci USA 94: 6238 (1997)). The fusion protein encoded by this plasmid was transiently expressed in 293-Epstein-Barr virus-encoded nuclear antigen (EBNA) cells and purified on a Protein A Sepharose immunoaffinity column using standard methods.

Six wells of a 96-well plate were coated overnight at 4 °C with goat anti-human IgG (Fc $\gamma$  fragment specific; Jackson Immunulogics) at a concentration of 25 $\mu$ g/ml in PBS (300 $\mu$ l/well). The wells were blocked for 1h at room temperature with 400 $\mu$ l of 1% BSA in PBS. After 3 washes with PBST (PBS + 0.05% Tween 20), 300 $\mu$ l hGFR $\alpha$ 3-Ig (10 $\mu$ g/ml in PBS containing 0.1% BSA) was added to each well. The plate was incubated for 1h at room temperature and shaken gently (200 strokes/min) to maximize the binding. The wells were then emptied and washed again 3 times with PBST. 250 $\mu$ l of conditioned media from a negative control CHO cell line or from the neublastin producing CHO cell line #25 was added to each of 3 wells. The plate was incubated for 3h at room temperature and shaken gently (300 strokes/min). The wells were then washed twice with PBST. 25 $\mu$ l of non-reducing Laemli loading buffer was added to the first well and the plate was shaken rapidly for 5 min to elute the bound proteins (1300 strokes/min). The content was transferred to the next well and the procedure was repeated to elute the proteins bound in the second and third wells. After adding



$\beta$ -mercaptoethanol (5% final), the samples were boiled for 5 minutes and analyzed by SDS-PAGE on a 10-20% polyacrylamide gel.

For western blot analysis, the proteins were transferred to nitrocellulose. The membrane was blocked and probed in 5% non fat dry milk, PBST and washed in PBST. Neublastin was detected by electrochemoluminescence after reaction with polyclonal antibodies (R30 and R31) raised against two neublastin peptides (at 1ug/ml) followed by reaction with HRP-conjugated goat anti-rabbit antibodies (BioRad). As shown in Figure 13, five neublastin specific bands were detected in the extracted proteins from the neublastin producing CHO cell line (lane 2). The lower two bands are very similar to the bands observed in Figure 12; again, the lower band probably represents the mature domain of neublastin generated after cleavage of the pro-domain.

Subsequent analysis (data not shown) of the bands in Figure 13 shows that deglycosylation with PGNase F of the approximately 18 kD band reduces that band to a size equivalent to the lower-most band in the gel of Figure 13. This suggests that neublastin may be produced as a glycosylated protein in mammalian cells.

#### Expression of Neublastin in *E. coli*

In order to express the neublastin gene in *E. coli*, syngenes were constructed with lower GC content and preferred *E. coli* codons. The syngene is being cloned into two vectors, pET19b and pMJB164, a derivative of pET19b. The construction with pET19b is shown in Figure 14. In this construct, the sequence encoding NBN113 is directly fused to an initiating methionine. Additional synthetic gene constructs for NBN are shown in SEQ ID NOS:52-54. The construction with pMJB164 is shown in Figure 15. In this construct, NBN113 is fused to a histidine tag (*i.e.* 10 histidines) separated by an enterokinase cleavage site. Additional synthetic gene constructs for NBN fused to a histidine tag are shown in SEQ ID NOS:55-57. The initiating methionine precedes the histidine tag.

#### Nucleotide sequence encoding neublastin in Figure 14

ATGGCTGGAGGACCGGGATCTCGTGCTCGTGCGAGCAGGAGCACGTGGCTGCTGCGTTCTCAACTA  
GTGCCGGTGCGTGCACTCGGACTGGGACACCGTTCCGACGAACTAGTACGTTTTCGTTTTGTTTCAGGA  
TCTTGTCGTGCGTGACGTTCTCCGCATGATCTATCTCTAGCATCTCTACTAGGAGCCGGAGCACTAAGA  
CCGCCGCCGGGATCTAGACCTGTATCTCAACCTTGTTGTAGACCTACTAGATACGAAGCAGTATCTTTC  
ATGGACGTAACTCTACATGGAGAACCGTAGATAGACTATCTGCAACCGCATGTGGCTGTCTAGGATGA

TAATAG (SEQ ID NO:29)

**Nucleotide sequence encoding his-tagged neublastin in Figure 15**

ATGGGCCATCATCATCATCATCATCATCATCACTCGAGCGGCCATATCGACGACGACACAAGGCT  
 5 GGAGGACCGGGATCTCGTGCTCGTGACAGGAGCACGTGGCTGTCTGCGTTCTCAACTAGTGCCG  
 GTGCGTGCACTCGGACTGGGACACCGTTCCGACGAACCTAGTACGTTTTTCGTTTTTGTTCAGGATCTTGT  
 CGTCGTGCACGTTCTCCGCATGATCTATCTCTAGCATCTCTACTAGGAGCCGGAGCACTAAGACCGCCG  
 CCGGGATCTAGACCTGTATCTCAACCTTGTGTAGACCTACTAGATACGAAGCAGTATCTTTCATGGAC  
 GTAAACTCTACATGGAGAACCGTAGATAGACTATCTGCAACCGCATGTGGCTGTCTAGGATGATAATAG  
 10 (SEQ ID NO:30)

**Example 6: Effect of Neublastin on the survival of Rat Embryonic Dopaminergic Neurons and ChAT Activity.**

15 In this series of experiments the effect of conditioned medium from neublastin-producing HiB5pUbi1zNBN22 cells described above was assessed.

**Preparation of Cultures:** The ventral mesencephalon or spinal cord was dissected out from rat E14 embryos in cold Hanks Buffered Salt Solution (HBSS). Tissue pieces were incubated in sterile filtered 0.1% trypsin (Worthington) and 0.05% DNase (Sigma) in HBSS at  
 20 37°C for 20 min. Tissue pieces was then rinsed four times in HBSS + 0.05% DNase and dissociated using a 1 ml automatic pipette. The suspension was then centrifuged at 600 rpm for 5 min and the pellet was re-suspended in serum conditioned medium (SCM; DMEM with 10% foetal calf serum). The total number of cells was assessed by trypan blue dye exclusion method and plated at a density of 100.000 cells/cm<sup>2</sup> in poly-L-lysine coated eight-well  
 25 chamber slides (Nunc) for assessment of dopaminergic neuron survival or at 200 000 cells/cm<sup>2</sup> in 48 well plates (Nunc) for ChAT activity measurements. Cells were incubated in SCM at 5% CO<sub>2</sub>/95% O<sub>2</sub> and 95% humidity in 37°C for 24-48h before changing to serum free medium (SFM) with addition of neurotrophic factors.

Cells for assessing dopaminergic neuron survival were left for 5 days in SFM +  
 30 trophic factor additions and then fixed for 5 min in 4% PFA and stained for tyrosine hydroxylase by immunohistochemistry.

Cells for ChAT activity were left for 3 days with SFM and then lysed in HBSS + 0.1% Triton X-100 and immediately frozen down on dry ice until Chat activity measurement.

**Trophic Factor Addition:** Conditioned medium was collected from non-transfected HiB5 control or HiB5 producing neublastin (HiB5pUbi1zNBN22) or GDNF (HiB5pUbi1zGDNF-L17). HiB5pUbi1zNBN22 produces approximately 20 ng GDNF/24 hours/10<sup>5</sup> cells as determined by GDNF-ELISA on conditioned medium, collected from the cells. The respective cell lines were incubated overnight with DMEM + 1% FCS and the supernatant was taken off and stored at -20°C until use. The supernatants were diluted in 1:50 in SFM when added to the cells. Separate wells were treated with HiB5 control supernatant (1:50) + purified recombinant rat GDNF (from 0.03 – 10 ng/ml).

The results of these experiments are shown in FIG. 4. FIGS. 4A-4C are illustrations of the effect of neublastin, secreted from HiB5pUbi1zNBN22 cells, on the survival of cultured rat embryonic, dopaminergic, ventral mesencephalic neurons and ChAT activity in cholinergic cranial nerve motor neurons in serum-free medium as described *infra* in Example 5.1.

FIG. 4A is an illustration of the dose-response curve for recombinant GDNF on ChAT activity (dpm/hour) measured at DIV5 in serum-free cultures which were initially established from E14 ventral mesencephali (*i.e.*, HiB5; GDNF 0.03 ng/ml; GDNF 0.1 ng/ml; GDNF 0.3 ng/ml; GDNF 1 ng/ml; GDNF 10 ng/ml; GDNF 100 ng/ml).

FIG. 4B is an illustration of ChAT activity (dpm/hour) measured at DIV5 in serum-free cultures which were initially established from E14 ventral mesencephali. Diluted conditioned medium from either neublastin producing HiB5pUbi1zNBN22 cells (neublastin) or GDNF-producing HiB5GDNFL-17 (GDNFL-17) cells were added as indicated in the figure (*i.e.*, neublastin 1:10; neublastin 1:50; GDNF L-17 1:50).

FIG. 4C is an illustration of the number of tyrosine hydroxylase immunoreactive cells per well (No. TH+ cells/well) at DIV7 in serum-free cultures which were initially established from E14 rat ventral mesencephali. Diluted conditioned medium from either non-transfected HiB5 cells (HiB5) or neublastin-producing HiB5pUbi1zNBN22 cells (neublastin) or recombinant GDNF, in various concentrations, were added as indicated in the figure (*i.e.*, HiB5 1:10; HiB5 1:40; GDNF 0.1 ng/ml; GDNF 10 ng/ml; GDNF 100 ng/ml; and neublastin 1:40).

Conditioned medium from neublastin transfected HiB5 cells diluted 1:40 significantly increases the number of TH immunoreactive cells per well compared to control (untransfected) HiB5 cells at an equivalent and a lower dilution (1:10 and 1:40) (see, *e.g.*, FIG. 4B). The increase in TH-immunoreactive cells is comparable to the increase seen at a maximal GDNF concentration (10 ng/ml). This indicates that neublastin secreted to the medium has an effect on

survival of the dopaminergic neuron population from rat embryonic ventral mesencephalon. In contrast, unlike GDNF secreted from transfected HiB5 cells, no effect of conditioned medium from neublastin transfected HiB5 cells is seen on another neuronal population in the same culture, the cholinergic neurons (see, e.g., FIG. 4A).

5

**Example 7: Effect of Neublastin on the survival of slice cultures of pig embryonic dopaminergic ventral mesencephalic neurons**

This experiment assessed the effect of co-culturing neublastin-producing HiB5pUbi1zNBN22 cells with slice cultures of ventral mesencephali from porcine embryos.

10 **Preparation of Cultures:** Ventral mesencephali (VM) were isolated from porcine embryos (E28; n=12) under sterile conditions, chopped into 400  $\mu$ m slices and placed in chilled Gey's balanced salt solution (GIBCO) with glucose (6.5 mg/ml). The tissue slices were cultured by the interface culture method, originally developed by *Stoppini et al.* (*L. Stoppini, P.A. Buchs, D. Muller, J. Neurosci. Methods* 1991 37 173-182).

15 In brief, slices were placed on semi-porous membranes (Millipore, 0.3  $\mu$ m; 8 slices/membrane corresponding to one VM) placed as inserts in 6-well plates (Costar) with serum containing medium (Gibco BRL). Each well contained 1 ml medium (50% Optimem, 25% horse serum, 25% Hank's balanced salt solution (all GIBCO)) supplemented with D-glucose to a final concentration of 25 mM. At day 0, 7000 transfected HiB5pUbi1zNBN22  
20 (neublastin) or 7000 non-transfected HiB5 cells (control) were seeded on each tissue slice. The co-cultures were first grown in an incubator at 33°C for 48 hours allowing the HiB5 cells immortalized with a temperature sensitive oncogene to proliferate, and then placed in an incubator at 37°C, where the HiB5 cells differentiate. The medium was changed twice a week. Antimitotics and antibiotics were not used at any stage.

25 **Determination of Dopamine by HPLC:** At day 12 and 21 *in vitro*, the culture medium was collected and analysed for dopamine using HPLC with electrochemical detection (*W.N. Slooth, J.B.P. Gramsbergen, J. Neurosci. Meth.* 1995 60 141-49).

**Tissue Processing and Immunohistochemistry:** At day 21, the cultures were fixed in 4% paraformaldehyde in phosphate buffer for 60 min., dehydrated in a 20% sucrose solution  
30 for 24 hours, frozen, cryostat sectioned at 20 $\mu$ m (4 series), and mounted onto gelatine coated microscope slides. One series of sections was immunostained for tyrosine hydroxylase (TH). Briefly, sections were washed in 0.05M tris-buffered saline (TBS, pH 7.4) containing 1%

Triton X-100 for 3x15 min. and incubated with 10% fetal bovine serum (FBS, Life Technologies) in TBS for 30 min. The tissue was then incubated for 24 hours at 4°C with monoclonal mouse anti-TH antibody (Boehringer Mannheim) diluted 1:600 in TBS with 10% FBS. After rinsing in TBS with 1% Triton X-100 for 3x15 min., sections were incubated for 60 min. with biotinylated anti-mouse IgG antibody (Amersham) diluted 1:200 in TBS with 10% FBS. The sections were then washed in TBS with 1% Triton X-100 (3x15 min.) and incubated for 60 min. with streptavidin-peroxidase (Dako) diluted 1:200 in TBS with 10% FBS. After washing in TBS (3x15 min.), bound antibody was visualised by treatment with 0.05% 3,3-diaminobenzidine (Sigma) in TBS containing 0.01% H<sub>2</sub>O<sub>2</sub>. Finally, the sections were dehydrated in alcohol, cleared in xylene, and cover-slipped in Eukitt.

**Cell counts and morphometric analysis:** Quantification of immunoreactive TH-ir neurons was performed using bright field microscopy (Olympus). Only cells displaying an intense staining with a well preserved cellular structure and a distinct nucleus were counted. The estimation was based on cell counts in every fourth culture section using a x20 objective. Cell numbers were corrected for double counting according to Abercrombie's formula (*M. Abercrombie, Anat. Rec.* 1946 **94** 239-47), using the average diameter of the nuclei in the TH-ir neurons ( $6.6 \pm 0.2 \mu\text{m}$ ,  $n = 30$ ). The size of the nuclei was estimated using a neuron tracing system (Neurolucida, MicroBrightField, Inc.).

The results of these experiments are shown in FIG. 5. FIGS. 5A-5C are illustrations of the effect of neublastin secreted from HiB5pUbi1zNBN22 cells on the function and survival of slice cultures of pig embryonic dopaminergic ventral mesencephalic neurons co-cultured with either HiB5pUbi1zNBN22 cells (neublastin) or HiB5 cells (control) as described *infra*.

FIG. 5A and FIG. 5B: illustrate dopamine released to the medium at DIV12 (Dopamine (pmol/ml) - day 12) and DIV21 (Dopamine (pmol/ml) - day 21), respectively. FIG. 5C is an illustration of the number of tyrosine hydroxylase immunoreactive cells per culture (TH-ir cells per culture) at DIV21.

At day 12 HPLC analysis revealed that medium from HiB5-neublastin co-cultures contained 84% more dopamine than medium from HiB5-C co-cultures (Fig. 5A). At day 21 the difference was 78% (Fig. 5B), and cell counts showed that HiB5-neublastin co-cultures contained 66% more tyrosine hydroxylase immunoreactive neurons than HiB5-C co-cultures ( $P < 0.05$ ) (Fig. 5C). This indicates that neublastin secreted from the HiB5pUbi1zNBN22 clone has a potent survival effect on embryonic porcine dopaminergic neurons.

**Example 8: Survival of Dorsal Root Ganglion Cells in Serum-free Medium**

This example shows the neurotrophic activity of a neublastin polypeptide in comparison with known neurotrophic factors.

5 Pregnant female mice were killed by cervical dislocation. The embryos were processed for culture as follows.

Electrolytically sharpened tungsten needles were used to dissect dorsal root ganglia from indicated stages of C57/Bl6 mice (Møllegaard Breeding, Denmark). Embryonic ganglia were incubated for 5 minutes at 37°C with 0.05% trypsin (Gibco/BRL) in calcium and  
 10 magnesium-free Hanks balanced salt solution. Postnatal ganglia were treated with collagenase/dispase 1mg/ml for 30 to 45 minutes and then trypsin/DNase 0,25% for 15 minutes. After removal of the trypsin solution, the ganglia were washed once with 10 ml of DMEM containing 10% heat inactivated horse serum, and were gently triturated with a fire-polished Pasteur pipette to give a single cell suspension.

15 The cells were plated on 24 well plates (Nunc), that were precoated with polyornithine (0.5 mg/ml, overnight) and laminin (20 mg/ml for 4 h; Gibco/BRL). The neurons were incubated at 37°C in a humidified 5% CO<sub>2</sub> incubator in a defined medium consisting of Hams F14 supplemented with 2 mM glutamine, 0.35% bovine serum albumin, 60 ng/ml progesterone, 16 mg/ml putrescine, 400 ng/ml L-thyroxine, 38 ng/ml sodium selenite, 340  
 20 ng/ml triiodo-thyronine, 60 mg/ml penicillin and 100 mg/ml streptomycin.

After 48 hours of incubation, neurons were clearly recognized by their bipolar morphology under phase-contrast optics. The percentage neuronal survival in the absence or presence of trophic factors (added to the culture medium prior to plating the neurons at 10 ng/ml), or of conditioned medium from the neublastin producing HiB5pUbi1zNBN22 cells,  
 25 was assessed by counting the neurons in the wells at 48 hours.

The results of these experiments are presented in FIG. 9, in which figure:

0 represents the control experiment (in absence of factors);

1 represents experiments in the presence of GDNF;

2 represents experiments in the presence of Neurturin;

30 3 represents experiments in the presence of Neublastin of the invention;

FIG. 9 presents data from experiments carried out on DRG cells isolated from developing fetuses at the various time points designated as follows:

E12 represents embryonic day 12;

E16 represents embryonic day 16;

P0 represents the day of birth;

P7 represents day 7 after birth; and

5 P15 represents day 15 after birth.

These results clearly show that the neurotrophic factor of the invention shows activities comparable to, or even better than those of, the well established neurotrophic factors.

#### **Example 9: *In vivo* effects of neublastin on nigral dopamine neurons**

10 In order to test the ability of neublastin (neublastin) to protect adult nigral dopamine (DA) neurons from 6-hydroxydopamine induced degeneration, we employed a rat model of Parkinson's disease (*Sauer and Oertel, Neuroscience* 1994 59, 401-415) and lentiviral gene transfer of neublastin.

**Lentivirus production:** To generate a lentiviral transfer vector encoding neublastin, 15 pHR'-neublastin, a 1331 bp BamHI fragment from neublastin cDNA was subcloned in the BamHI/Bgl II site of pSL301 (Invitrogen). From this construct a 1519 bp BamHI/XhoI fragment was cut out and ligated in the BamHI/XhoI site of pHR' carrying a woodchuck hepatitis virus post-translational fragment (*Zufferey R, Donello JE, Trono D, Hope TJ: Woodchuck hepatitis virus posttranscriptional regulatory element enhances expression of* 20 *transgenes delivered by retroviral vectors*"; *J. Virol.* 1999 73 (4) 2886-2892). To generate pHR'-GDNF a 701 bp BamHI/XhoI fragment from pUbi1z-GDNF was ligated in the BamHI/XhoI site of pHR'.

Production of the lentiviral vector have been described by *e.g. Zufferey et al. (Zufferey R, Nagy D, Mandel RJ, Naldini L, Trono D: "Multiply attenuated lentiviral vector achieves* 25 *efficient gene delivery in vivo*"; *Nat. Biotechnol.* 1997 15 (9) 871-875). Briefly, the transfer constructs and the helper plasmids pR8.91 and pMDG were co-transfected into 293T cells. Virions released into the media were collected at 48 and 72 hrs post-transfection. To concentrate the virus, the media was centrifuged 1.5 hrs at 141 000g, and the pellet dissolved in DMEM. The titer of a control carrying the gene for Green Fluorescent Protein ("GFP") was 30 determined to be  $10^8$  transforming units (TU) /ml by GFP fluorescence in 293T cells. A RNA slot blot technique (*von Schwedler U, Song J, Aiken C, Trono D: "Vif is crucial for human immunodeficiency virus type 1 proviral DNA synthesis in infected cells*"; *J. Virol.* 1993 67 (8)

4945-4955) was used to determine viral particle titer. In the GDNF supernatant and neublastin supernatant there was 10 times less particles as compared to the GFP supernatant.

**Surgical Procedures:** All work involving animals was conducted according to the rules set by the Ethical Committee for Use of Laboratory Animals at Lund University.

5 A total of 21 young adult female Sprague-Dawley rats (B&K Universal, Stockholm, Sweden) were used and housed under 12 hours light:dark cycle with free access to rat chow and water. Retrograde labelling and 6-OHDA lesions were performed 3 weeks prior to lesion according to Sauer and Oertel [*Sauer and Oertel, Neuroscience* 1994 59:401-415]. Briefly, under Equithesin anaesthesia (0.3 ml/100g) the rats were injected bilaterally with 0.2 µl of a 2% solution (dissolved in 0.9% NaCl) of the retrograde tracer Fluoro-Gold (FG; Fluorochrome, Inc., Englewood, CO). Injections were made using a 2 µl Hamilton syringe at co-ordinates: AP= +0.5 mm; ML= ±3.4 mm relative to bregma; DV= -5.0 mm relative to the dura and incisor bar set to 0.0 mm. In addition, 0.05 µl/min was injected with another 5 min left before the needle was retracted.

15 Fourteen days after the FG injections animals received a total of 5 deposits (1 µl/deposit) of a lentiviral vector carrying the gene for green fluorescent protein (GFP), neublastin or GDNF. Four of the deposits were into the striatum along two needle tracts at the following co-ordinates: AP= +1.0 mm, ML= -2.6mm, DV<sub>1</sub>=-5.0mm DV<sub>2</sub>=-4.5mm and AP= 0.0mm, ML= -3.7mm, DV<sub>1</sub>=-5.0mm DV<sub>2</sub>=-4.5mm. The supranigral deposit was made at 20 AP= -5.2 mm, ML= -2.0 mm, DV<sub>1</sub>=-6.3 mm. Tooth bar was set at -2.3 mm.

Twenty-one days after retrograde labelling, and 7 days after lentiviral injections the animals were re-anaesthetised and with a 10 µl Hamilton syringe a single deposit of 20 µg 6-OHDA (Sigma; calculated as free base and dissolved in 3 µl ice cold saline supplemented with 0.02% ascorbic acid) was injected into the right striatum in the same location as the FG deposits. The injection rate was 1 µl/min, leaving another 3 min before retracting the needle.

**Tissue Processing:** At 21 days after the 6-OHDA injection the animals were deeply anaesthetised with chloral hydrate and transcardially perfused with saline (pH 7.4; room temperature) for one min followed by 200 ml ice cold formaldehyde solution (4% paraformaldehyde in 0.1M phosphate buffer, pH 7.4). The brains were dissected and postfixed 30 in the same fixative for 3-4 hours and then transferred into 25% sucrose/0.1M phosphate buffer for 48 hours. Five series of 40 µm sections through the striatum and substantia nigra (SN) were cut on a freezing microtome.



**Quantitative Assessment of Dopaminergic Neurons in the SN:** The number of FG-labelled in the SN pars compacta was assessed by a blinded observer as described previously [Sauer and Oertel, *Neuroscience* 1994 59, 401-415]. In brief, three consecutive sections centred around the level of the medial terminal nucleus of the accessory optic tract (MTN; -5.3

5 in the atlas of Paxinos and Watson (1997)) were used and all labelled/stained neurons laterally to the MTN was counted at 40 x magnification (n=6-7/group). FG-labelled neurons were included if they were brightly fluorescent under epi-illumination at 330 nm, displayed a neuronal profile and extend at least one neuritic process.

On the lesion side in animals receiving injections of lentivirus carrying GFP the number

10 of FG-positive nigral neurons were reduced to 18% of that on the intact side. In contrast, animals injected with lenti-neublastin showed a near complete protection of the number of FG-positive nigral neurons (89%). This was as efficient as lenti-GDNF treated animals where 87% of the retrogradely labelled neurons remained on the lesioned side. This shows that Neublastin is a potent survival factor for lesioned adult nigral dopamine neurons and that it is as potent as

15 GDNF.

FIG. 6 is an illustration of the *in vivo* effect of lentiviral-produced Neublastin on nigral dopamine neurons. Neurons of the SN pars compacta, in female Sprague Dawley rats, were retrogradely-labelled with Fluorogold (FG), 3 weeks prior to a single injection of 6-hydroxydopamine (6-OHDA) in the right striatum. One week before the 6-OHDA injection,

20 the animals received injections with lentiviral vectors expressing Neublastin (neublastin), GDNF (GDNF) or the Green Fluorescent Protein (GFP) as indicated in the figure. Twenty one days after the 6-OHDA injections, the number of FG-labelled neurons in both sides of the striata were determined. The figure shows the percentage (%FG lesion/intact) of FG-labelled neurons in the lesioned (right) side versus the intact (left) side of the striata of the three groups

25 of animals.

#### **Example 10: Production of Antibodies**

To prepare antibodies against Neublastin, two rabbits were immunised with either peptide 1: CRPTRYEAVSFMDVNST (amino acids 108-124 of SEQ ID NO:9); or peptide 2:

30 ALRPPPGSRPVSQPC (amino acids 93-107 of SEQ ID NO:9) conjugated to carrier protein at 3 week intervals. Two rabbits for each peptide were immunized at week 0, 3, 6 and 10, and

bleeds were collected at week 7 and 11. The second bleed was affinity purified via a peptide affinity column. The antibodies were named Ab-1 and Ab-2, according to the peptide.

Western blot: 2 x 10<sup>6</sup> HiB5 cells, stably transfected with the cDNA for neublastin (Hib5pUbi1zNBN22), or untransfected HiB5 cells, were incubated overnight in serum free medium with N<sub>2</sub> supplement (GIBCO). The medium was concentrated on small concentrators with cut-off membranes of 5 kDa (Millipore, Bedford, MA). Concentrated samples were added 5 x Laemmli sample buffer and were heated to 95°C for 5 minutes. Samples were separated by SDS polyacrylamide gel electrophoresis on 15% acrylamide gels and transferred to PVDF-membranes. Residual protein-binding sites were blocked with 5% non-fat dry milk in PBS with 0.1% Tween-20. Membranes were incubated overnight with neublastin antibody (1:1000), followed by incubation with a secondary anti-rabbit or anti-mouse IgG antibody conjugated to horseradish peroxidase (1:2000).

Immunostaining was visualized using enhanced chemiluminescence Plus (ECL+) according to the manufacturer's instructions (Amersham). The results of these experiments are shown in FIG. 3 and Example 5.

Using standard techniques, we also raised rabbit polyclonal antibodies against the following peptides:

Peptide R27: GPGSRARAAGARGC	(amino acids 30-43 of SEQ ID NO:9);
Peptide R28: LGHRSDELVRFRFC	(amino acids 57-70 of SEQ ID NO:9);
Peptide R29: CRRARSPHDLSL	(amino acids 74-85 of SEQ ID NO:9);
Peptide R30: LRPPGSRPVSQPC	(amino acids 94-107 of SEQ ID NO:9); and
Peptide R31: STWRTVDRLSATAAC	(amino acids 123-136 of SEQ ID NO:9).

Only peptides R30 and R31, relatively close to the C-terminus, recognized the denatured protein under reducing conditions on a Western blot.

**Example 11: Biological Activity of a Truncated Rat Neublastin Polypeptide Comprising the Last Carboxy-terminal 102 Amino Acids.**

30

The amino acid sequence of rat prepro-neublastin is provided below as SEQ ID NO: 34:

1 MELGLGEPTA LSHCLRPRWQ PALWPTLAAL ALLSSVTEAS LDPMSRSPAS

51 RDVPSPVLAP PTDYLPGGHT AHLCSERALR PPPQSPQPAP PPPGPALQSP  
 113aa N-7 -9 -11 -14  
 101 PAALRGARAA RAGTRSSRAR ATDARGCRLR SQVVPVSALG LGHSSDELIR  
 151 FRFCSGSCRR ARSPHDLSLA SLLGAGALRS PPGSRPISQP CCRPTRYEAV  
 5 201 SFMDVNSTWR TVDHLSATAC GCLG (SEQ ID NO: 34)

Each of the neublastin truncations described below is identified above the corresponding starting residue (bold letters in the above sequence). The start of the 113 amino acid neublastin form (NBN113) is also labeled.

- 10 The biological activity of a truncated form of neublastin containing the 102 carboxy terminal amino acids of rat neublastin was examined in cells. Truncated neublastin was generated by digesting a 113 amino acid form of rat neublastin, having the amino acid sequence containing amino acids 111-224 (rat NBN113, underlined above) of SEQ ID NO: 34, with a non-specific aminopeptidase (Sigma, MO) for two hours at room temperature.
- 15 Following this digestion, neublastin was subjected to size exclusion chromatography (SEC) to separate any contaminating components from the protein. The molecular weight of truncated neublastin (designated "NBN113 (N-11)") was confirmed by mass spectroscopy and determined to be 10.931 kDa, the predicted weight of the 102 amino acid polypeptide (NBN102). As a control, rat NBN113 was treated in parallel with enzyme buffer (25 mM Tris
- 20 pH 8.) alone. No digestion was observed with this control, and the proper 12.047 kDa polypeptide was identified.

- The biological activities of NBN113, NBN113 treated with enzyme buffer and NBN113 (N-11) were compared in a cellular RET (*c-RET*) activation assay. Neublastin activity was determined by its ability to stimulate *c-RET* phosphorylation in NB41A3-mRL3 cells, an
- 25 adherent murine neuroblastoma cell line stably transfected with murine GFR $\alpha$ 3 and that expresses RET and GFR $\alpha$ 3. NB41A3-mRL3 cells were plated in DMEM supplemented with 10 % FBS at  $2 \times 10^5$  cells per well in 24-well plates, and cultured for 18 hours at 37°C and 5 % CO<sub>2</sub>. Following removal of the media and a cell wash with 1 ml of PBS per well, the cells were stimulated with DMEM containing either NBN113, NBN113 treated with enzyme buffer,
- 30 or NBN113 (N-11) for 10 minutes at 37°C and 5 % CO<sub>2</sub>. To stop activity, the media was removed and the cells washed with PBS immediately before lysis with 10 mM Tris, pH 8.0, 0.5 % NP40, 0.2 % DOC, 50 mM NaF, 0.1 mM Na<sub>3</sub>VO<sub>4</sub>, and 1 mM PMSF. After a 1-hour incubation at 4°C, the lysates were agitated by repeated pipetting and transferred (0.25 ml per well) to a 96-well ELISA plate coated with anti-RET mAb (AA.GE7.3). The wells were

blocked at room temperature for 1 hour with blocking buffer (TBST containing 1 % normal mouse serum and 3 % BSA in TBS buffer (15 mM Tris pH 7.4, 150 mMNaCl)) followed by six washes with TBST (TBS + 0.05 % Tween 20) alone.

Phosphorylated RET was detected by incubating (2 hours) the captured receptor with  
5 4G10 (Upstate Biotechnology, NY) an HRP-conjugated phosphotyrosine antibody (0.2 ug per well). Following the incubation, the wells were washed six times with TBST, and the HRP activity detected at 450 nm with a colorimetric assay. The absorbance values from wells treated with lysate or with lysis buffer alone were measured, background corrected, and the data plotted as a function of the concentration of neublastin present in the activation mixture. The  
10 results are shown in FIG. 16. The absorbance values from wells treated with lysate or with lysis buffer were measured and the background-corrected signals were plotted as a function of the concentration of neublastin.

These results demonstrate that the amino-terminal truncated form (102 amino acids) of neublastin exhibits cellular biological activity that is indistinguishable from that of the  
15 NBN113 form of neublastin.

**Example 12: Biological Activity of Three Amino-terminal Truncated Forms (99, 104, and 106 amino acids) of Neublastin.**

The activities of truncated neublastin polypeptides (99 and 106 amino acids) in the  
20 cellular RET activation assay were examined, and the results shown in FIG. 17. In addition to the 99 and 106 amino acid forms, two other neublastin molecules were included in this assay for comparison: Rat NBN113, which served as a reference, and a 113 amino acid rat neublastin mutein in which the arginine at position 14 of the rat NBN113 sequence was replaced with a lysine residue ("NBN R14K (N)"). This mutein facilitated the generation of the 99 amino acid  
25 form, lacking the fourteen amino terminal residues ("NBN R14K (N-14)"), when cleaved at this site using a lysine-specific protease (Endo Lys C; WAKO, VA). The 106 amino acid form, lacking the seven amino terminal residues ("NBN 113 (N-7)"), was generated from rat NBN113 by partial proteolysis with trypsin (Biozyme, San Diego). All truncated products were characterized by mass spectroscopy as above in Example 11.

30 The activities of NBN113, NBN R14K (N), NBN R14K (N-14), and NBN113 (N-7) were examined at concentrations from 0.005 nM to 100 nM, 0.05 nM to 100 nM, 0.006 nM to 114 nM and 0.05 nM to 107 nM, respectively. Activity was measured as absorbance at  $A_{450nm}$

as described in Example 11. Each neublastin form displayed similar activity at each concentration tested. These results demonstrate that truncated forms of rat NBN113 lacking either the 14 or 7 amino-terminal residues are as active as rat NBN113 in the KIRA activation assay.

5           The KIRA activation assay shown in FIG. 18 was performed to determine the biological activity of a 104 amino acid neublastin form ("NBN104 (N-9)"). This polypeptide, lacking nine amino-terminal residues present in the wild-type rat NBN113 molecule, was generated by treating CHO-expressed rat NBN113 with trypsin. As before, the molecular mass was confirmed by mass spectroscopy. The activity of the two polypeptides (NBN104 (N-9) and  
10 NBN113) was examined at concentrations from 0.05 nM to 115 nM. Activity was measured as absorbance at A<sub>450nm</sub> as described in Example 11. Activity of the N-9 form was at least as high as the activity of the rat NBN113. These results demonstrate that a truncated form (104 amino acids) lacking the 9 amino terminal amino acids of the rat NBN113 polypeptide is at least as active as rat NBN113 in the KIRA ELISA activation assay.

15           Table 5 illustrates the relationship between the disclosed prepro neublastin polypeptide sequences of the invention. Line 1 provides the polypeptide of SEQ ID NO:2, line 2 provides the polypeptide of SEQ ID NO:4 and line 3 provides the polypeptide of SEQ ID NO:9. The seven conserved cysteine residues are designated by symbols ("\*", "#", "+" and "|") to indicate the intramolecular (\* with \*, # with #, and + with +) and intermolecular ("|") disulfide bridges  
20 formed in the mature dimerized neublastin ligand. The amino terminus of each truncated neublastin polypeptides is designated by "o" for NBN112 through NBN99, respectively.

Table 5 Alignment of Neublastin Polypeptides									
	10	20	30	40	50				
SEQ: 2	..... ..... ..... ..... ..... ..... ..... ..... ..... .....				MPALWPTLAALALL	14			
SEQ: 4	MPGLISARGQPL	LEVLPPQAH	LGALFLPEAPLGLSA	QPALWPTLAALALL		50			
SEQ: 9	-----M	ELG-----	LGGLSTLSHCPWPRR	QPALWPTLAALALL		33			
	60	70	80	90	100				
SEQ: 2	SSVAEASLGSA	PRSPAPREG	PPFVLAS	PAGHLPGGRTAR	WCSGRARRRR	64			
SEQ: 4	SSVAEASLGSA	PRSPAPREG	PPFVLAS	PAGHLPGGRTAR	WCSGRARRPPP	100			
SEQ: 9	SSVAEASLGSA	PRSPAPREG	PPFVLAS	PAGHLPGGRTAR	WCSGRARRPPP	83			
	110	120	130	140	150				
SEQ: 2	RHFSARAPAACT	PTICSSPR-V	RAARLGCRA	ARS	SGSCA-GCRLRSOLV	112			

SEQ: 4	-QPSRPAPPPAPPSALPRGGRAARAGGPGN	RARAAGARGCRLRSOLVPV	149	
SEQ: 9	-QPSRPAPPPAPPSALPRGGRAARAGGPGS	RARAAGARGCRLRSOLVPV	132	
	oooooooooooooooo*			
	160	170	180	
	190	200		
SEQ: 2	RALGLGHRDELVRFRFC	TGSCPRARSPHDL	SLASLLGAGALRPPPGSRP	162
SEQ: 4	RALGLGHRDELVRFRFC	SGSCRRARSPHDL	SLASLLGAGALRPPPGSRP	199
SEQ: 9	RALGLGHRDELVRFRFC	SGSCRRARSPHDL	SLASLLGAGALRPPPGSRP	182
		#	+	
	210	220	230	
	..... .....			

### EQUIVALENTS

From the foregoing detailed description of the specific embodiments of the invention, it should be apparent that particular novel compositions and methods involving nucleic acids, polypeptides, antibodies, detection and treatment have been described. Although these particular embodiments have been disclosed herein in detail, this has been done by way of example for purposes of illustration only, and is not intended to be limiting with respect to the scope of the appended claims that follow. In particular, it is contemplated by the inventors that various substitutions, alterations, and modifications may be made as a matter of routine for a person of ordinary skill in the art to the invention without departing from the spirit and scope of the invention as defined by the claims. Indeed, various modifications of the invention in addition to those described herein will become apparent to those skilled in the art from the foregoing description and accompanying figures. Such modifications are intended to fall within the scope of the appended claims.

### Description of Sequences Contained in the Sequence Listing

SEQ ID NO:1	Human neublastin nucleic acid.	865 bp
SEQ ID NO:2	Human neublastin polypeptide from sequence 1.	200 aa
SEQ ID NO:3	Coding region (CDS) of a human pre-pro- polypeptide.	861 bp
SEQ ID NO:4	Human neublastin polypeptide from sequence 3.	238 aa
SEQ ID NO:5	Variant of human neublastin in sequence 4 (Xaa1 is Asn or <u>Thr</u> ; Xaa2 is <u>Ala</u> or Pro).	140 aa
SEQ ID NO:6	Variant of human neublastin in sequence 4 (Xaa1 is Asn or <u>Thr</u> ; Xaa2 is <u>Ala</u> or Pro).	116 aa
SEQ ID NO:7	Variant of human neublastin in sequence 4 (Xaa1 is Asn or <u>Thr</u> ; Xaa2 is <u>Ala</u> or Pro).	113 aa

	SEQ ID NO:8	cDNA from positive colony PCR of human fetal brain cDNA.	861 bp
	SEQ ID NO:9	human fetal brain pre-pro-neublastin polypeptide including "stop" (corresponds to seq. 8)	221 aa
	SEQ ID NO:10	Variant of pre-pro-neublastin (seq. 9) NBN140, 14.7 kD.	140 aa
5	SEQ ID NO:11	Variant of pre-pro-neublastin (seq. 9) NBN116, 12.4 kD.	116 aa
	SEQ ID NO:12	Variant of pre-pro-neublastin (seq. 9) NBN113, 12.1 kD.	113 aa
	SEQ ID NO:13	PCR product from screen of human fetal brain cDNA master plate using SEQ ID NOS:17 and 18 as primers.	102 bp
10	SEQ ID NO:14	PCR product from screen of mouse fetal cDNA master plate using SEQ ID NOS:21 and 22 as primers.	220 bp
	SEQ ID NO:15	Full length mouse Neublastin cDNA.	2136 bp
	SEQ ID NO:16	Mouse pre-pro-neublastin polypeptide.	224 aa
	SEQ ID NO:17	"NBNint.sense" Top Primer for NBN from human fetal brain cDNA complementary to bases 551-568 of SEQ ID NO:1	18 nt
15	SEQ ID NO:18	"NBNint.antisense" Bottom Primer for NBN from human fetal brain cDNA reverse complement to bases 633-652 of SEQ ID NO:1	20 nt
	SEQ ID NO:19	"NBNnext.sense" Top Primer for whole human brain mRNA RT-PCR complementary to bases 58-74 of SEQ ID NO:8.	17 nt
20	SEQ ID NO:20	"NBNnext.antisense" Bottom Primer for whole human brain mRNA RT-PCR reverse complement to bases 850-865 of SEQ ID NO:8.	16 nt
	SEQ ID NO:21	"NBNint.sense" NBN C2 Primer for screening mouse fetal cDNA master plate complementary to bases 1398-1415 of SEQ ID NO:15.	18 nt
	SEQ ID NO:22	"NBNint.antisense" NBN C2as Primer for screening mouse fetal cDNA master plate. Reverse complement to bases 1598-1617 of SEQ ID NO:15.	20 nt
25	SEQ ID NO:23	Primer Pair 1 Sense PCR Primer for human genomic DNA amplification complementary to bases 60-88 of SEQ ID NO:3.	29 nt
	SEQ ID NO:24	Primer Pair 1 Antisense PCR Primer for human genomic DNA amplification Reverse complement to bases 835-861 of SEQ ID NO:3.	27 nt
30	SEQ ID NO:25	Primer Pair 2 Sense PCR Primer for human genomic DNA amplification complementary to bases 1-35 of SEQ ID NO:3.	35 nt
	SEQ ID NO:26	Primer Pair 2 Antisense PCR Primer for human genomic DNA amplification reverse complement to bases 786-819 of SEQ ID NO:3.	34 nt
	SEQ ID NO:27	Antisense alkaline phosphatase conjugated hybridization probe, complementary to bases 1140-1169 of mouse neuroblastin cDNA.	30 nt
35	SEQ ID NO:28	"NBNnext.sense" Top Primer for whole human brain mRNA RT-PCR complementary to bases 1-16 of SEQ ID NO:1	16 nt
	SEQ ID NO:29	Syngene ORF from Figure 14 of Neublastin.	351 nt
	SEQ ID NO:30	Syngene ORF from Figure 15 of HisNeublastin.	414 nt
	SEQ ID NO:31	Primer for isolating Neublastin.	39 nt
40	SEQ ID NO:32	Primer for isolating Neublastin.	39 nt
	SEQ ID NO:33	"NBNint.antisense" NBN primer; reverse complement to bases 715-730 of SEQ ID NO:8.	16 nt
	SEQ ID NO:34	Rat pre-pro-neublastin	224 aa
	SEQ ID NO:35	Human Neublastin (NBN112)	112 aa
45	SEQ ID NO:36	Human Neublastin (NBN111)	111 aa
	SEQ ID NO:37	Human Neublastin (NBN110)	110 aa
	SEQ ID NO:38	Human Neublastin (NBN109)	109 aa
	SEQ ID NO:39	Human Neublastin (NBN108)	108 aa

	SEQ ID NO:40	Human neublastin (NBN107)	107 aa
	SEQ ID NO:41	Human neublastin (NBN106, N-7)	106 aa
	SEQ ID NO:42	Human neublastin (NBN105)	105 aa
	SEQ ID NO:43	Human neublastin (NBN104, N-9)	104 aa
5	SEQ ID NO:44	Human neublastin (NBN103)	103 aa
	SEQ ID NO:45	Human neublastin (NBN102)	102 aa
	SEQ ID NO:46	Human neublastin (NBN101)	101 aa
	SEQ ID NO:47	Human neublastin (NBN100)	100 aa
	SEQ ID NO:48	Human neublastin (NBN99, N-14)	99 aa
10	SEQ ID NO:49	Neurturin - Table 3	197 aa
	SEQ ID NO:50	Persephin - Table 3	156 aa
	SEQ ID NO:51	GDNF - Table 3	211 aa
	SEQ ID NO:52	Synthetic Gene for Neublastin	365 nt
	SEQ ID NO:53	Synthetic Gene for Neublastin	365 nt
15	SEQ ID NO:54	Synthetic Neublastin	114 aa
	SEQ ID NO:55	Synthetic Gene for HisNeublastin	442 nt
	SEQ ID NO:56	Synthetic Gene for HisNeublastin	442 nt
	SEQ ID NO:57	Synthetic HisNeublastin	135 aa



## CLAIMS

We Claim:

1. A truncated neublastin polypeptide, wherein the amino terminus of said truncated neublastin polypeptide lacks one or more amino-terminal amino acids of a mature neublastin polypeptide.
2. The truncated neublastin polypeptide of claim 1, wherein said truncated neublastin polypeptide, when dimerized, binds to a RET polypeptide.
3. The truncated neublastin polypeptide of claim 1, wherein said truncated neublastin polypeptide, when dimerized, induces dimerization of said RET polypeptide.
4. The truncated neublastin polypeptide of claim 1, wherein said truncated neublastin polypeptide includes seven cysteine residues at positions corresponding to positions 16, 43, 47, 80, 81, 109, and 111 of the neublastin polypeptide sequence of SEQ ID NO:12.
5. A polypeptide comprising the amino acid sequence of a truncated neublastin polypeptide, wherein the amino acid sequence of said truncated neublastin polypeptide is less than 113 amino acids in length and includes an amino acid sequence at least 70% homologous to amino acids 42-140 of SEQ ID NO:9.
6. The polypeptide of claim 5, wherein the amino acid sequence of said truncated neublastin polypeptide is at least 80% homologous to amino acids 42-140 of SEQ ID NO:9.
7. The polypeptide of claim 5, wherein the amino acid sequence of said neublastin polypeptide is at least 90% homologous to amino acids 42-140 of SEQ ID NO:9.
8. The polypeptide of claim 5, wherein the amino acid sequence of said neublastin polypeptide is at least 95% homologous to amino acids 42-140 of SEQ ID NO:9.

9. The polypeptide of claim 5, wherein the amino acid sequence of said truncated neublastin polypeptide comprises amino acids 42-140 of SEQ ID NO:9.
10. The polypeptide of claim 5, wherein the amino acid sequence of said truncated neublastin polypeptide consists essentially of 99 amino acids.
11. The polypeptide of claim 5, wherein the amino acid sequence of said truncated neublastin polypeptide is at least 80% homologous to amino acids 39-140 of SEQ ID NO:9.
12. The polypeptide of claim 5, wherein the amino acid sequence of said neublastin polypeptide is at least 90% homologous to amino acids 39-140 of SEQ ID NO:9.
13. The polypeptide of claim 5, wherein the amino acid sequence of said neublastin polypeptide is at least 95% homologous to amino acids 39-140 of SEQ ID NO:9.
14. The polypeptide of claim 5, wherein the amino acid sequence of said truncated neublastin polypeptide comprises amino acids 39-140 of SEQ ID NO:9.
15. The polypeptide of claim 5, wherein the amino acid sequence of said truncated neublastin polypeptide consists essentially of 102 amino acids.
16. The polypeptide of claim 5, wherein said truncated neublastin polypeptide is obtained by:  
providing a mature neublastin polypeptide; and  
contacting said mature neublastin polypeptide with at least one protease under conditions sufficient to produce said truncated neublastin polypeptide.
17. The polypeptide of claim 16, wherein said truncated neublastin polypeptide is produced as an exoprotease neublastin polypeptide digestion product by contacting said mature neublastin polypeptide with at least one exoprotease.
18. The polypeptide of claim 16, wherein said exoprotease is chosen from the group consisting of amino peptidase, Endo Lys C, and trypsin.

19. The polypeptide of claim 16, further comprising contacting said exopeptidase neublastin polypeptide digestion product with a dipeptidyl peptidase.
20. The polypeptide of claim 5, wherein said truncated neublastin polypeptide is glycosylated.
21. A nucleic acid comprising an open reading frame which encodes the polypeptide of claim 5.
22. A nucleic acid that hybridizes specifically under high stringency solution hybridization conditions to the nucleic acid of claim 21.
23. A method of using the neublastin nucleic acid of claim 21, comprising the step of causing a polypeptide encoded by said nucleic acid to be expressed in a cell.
24. The method of claim 23, further comprising the step of administering said nucleic acid to an animal, and causing said polypeptide to be expressed in said animal.
25. A vector comprising the truncated neublastin nucleic acid of claim 21.
26. The vector of claim 25, wherein said vector is an expression vector.
27. A method of using the vector of claim 26, comprising the step of causing a polypeptide encoded by said nucleic acid to be expressed from said nucleic acid.
28. A cell transformed with the nucleic acid of claim 21.
29. The cell of claim 28, wherein said cell is selected from the group consisting of a mammalian cell, a fungal cell, a yeast cell, an insect cell, and a bacterial cell.
30. The method of claim 29, wherein said mammalian cell is a Chinese hamster ovary cell.

31. The method of claim 29, wherein said mammalian cell is a cell derived from the mammalian central nervous system.
32. A method of making the truncated neublastin polypeptide of claim 5, said method comprising the step of expressing said polypeptide from the nucleic acid of claim 21.
33. The method of claim 32, comprising the step of culturing a cell comprising said nucleic acid in a culture medium which permits the production of said truncated neublastin polypeptide.
34. The method of claim 33, further comprising the step of recovering said polypeptide from said culture medium.
35. A purified truncated neublastin polypeptide obtained by the method of claim 32.
36. A pharmaceutical composition comprising a truncated neublastin polypeptide and a pharmaceutically acceptable carrier.
37. A pharmaceutical composition comprising a nucleic acid encoding a truncated neublastin polypeptide and a pharmaceutically acceptable carrier.
38. A method of administering the truncated neublastin polypeptide of claim 1, comprising the step of delivering said polypeptide to an isolated cell or *in vivo* to a mammal.
39. The method of claim 38, wherein said administration *in vivo* comprises systemic administration.
40. The method of claim 39, wherein said mammal is afflicted with a condition selected from the group consisting of ischemic neuronal damage, traumatic brain injury, peripheral neuropathy, neuropathic pain, Alzheimer's disease, Huntington's disease, Parkinson's disease, amyotrophic lateral sclerosis, and memory impairment.

41. The method of claim 39, wherein said mammal is afflicted with a neuronal disorder of the peripheral nervous system, the medulla, or the spinal cord.
42. A method of treating a neurodegenerative disease or disorder in an animal, comprising administering to said animal the truncated neublastin nucleic acid of claim 21.
43. A method of treating a neurodegenerative disease or disorder in an animal, comprising administering to said animal the truncated neublastin polypeptide of claim 1 or the polypeptide of claim 5.
44. A method of treating a peripheral neuropathy in a mammal, comprising administering a therapeutically effective amount of a truncated neublastin polypeptide to said mammal.
45. The method of claim 44, wherein said peripheral neuropathy is selected from the group consisting of trauma-induced neuropathies, viral-induced neuropathies, chemotherapy-induced neuropathies, toxin-induced neuropathies, drug-induced neuropathies, vitamin-deficiency-induced neuropathies; idiopathic neuropathies; and diabetic neuropathies.
46. The method of claim 44, wherein the truncated neublastin polypeptide is delivered directly into the central nervous system.
47. The method of claim 44, wherein the truncated neublastin polypeptide is delivered systemically by subcutaneous injection, intravenous administration, or intravenous infusion.
48. A method of treating neuropathic pain in a mammal, comprising administering a therapeutically effective amount of a truncated neublastin polypeptide to said mammal.
49. The method of claim 48, wherein said neuropathic pain is associated with toxin-induced nerve damage, pathogen-induced nerve damage, trauma-induced nerve damage, drug-induced nerve damage, idiopathic neuropathy, diabetic neuropathy, inflammation-induced nerve damage, or neurodegeneration.

50. A method of treating a peripheral neuropathy in a mammal, the method comprising administering a therapeutically effective amount of a nucleic acid encoding truncated neublastin polypeptide to said mammal.
51. The method of claim 50, wherein said peripheral neuropathy is selected from the group consisting of trauma-induced neuropathies, viral-induced neuropathies, chemotherapy-induced neuropathies, toxin-induced neuropathies, drug-induced neuropathies, vitamin-deficiency-induced neuropathies; idiopathic neuropathies; and diabetic neuropathies.
52. The method of claim 50, wherein the nucleic acid encoding said truncated neublastin polypeptide is delivered directly into the central nervous system.
53. The method of claim 50, wherein the truncated neublastin polypeptide is delivered systemically by subcutaneous injection, intravenous administration, or intravenous infusion.
54. A kit comprising, in one or more containers, a substance selected from the group consisting of a truncated neublastin polypeptide and a nucleic acid encoding a truncated neublastin polypeptide.
55. The polypeptide of claim 5, wherein the amino acid sequence of said truncated neublastin polypeptide comprises amino acids 41-140 of SEQ ID NO:9.
56. The polypeptide of claim 5, wherein the amino acid sequence of said truncated neublastin polypeptide consists essentially of 100 amino acids.
57. The polypeptide of claim 5, wherein the amino acid sequence of said truncated neublastin polypeptide comprises amino acids 40-140 of SEQ ID NO:9.
58. The polypeptide of claim 5, wherein the amino acid sequence of said truncated neublastin polypeptide consists essentially of 101 amino acids.

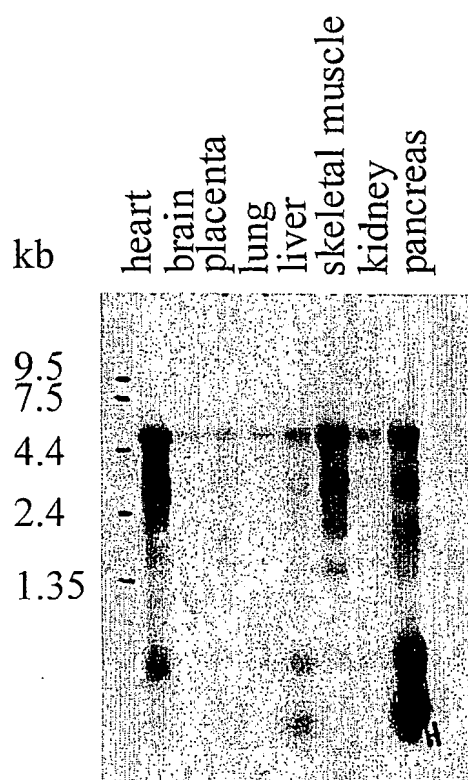
59. The polypeptide of claim 5, wherein the amino acid sequence of said truncated neublastin polypeptide comprises amino acids 38-140 of SEQ ID NO:9.
60. The polypeptide of claim 5, wherein the amino acid sequence of said truncated neublastin polypeptide consists essentially of 103 amino acids.
61. The polypeptide of claim 5, wherein the amino acid sequence of said truncated neublastin polypeptide comprises amino acids 37-140 of SEQ ID NO:9.
62. The polypeptide of claim 5, wherein the amino acid sequence of said truncated neublastin polypeptide consists essentially of 104 amino acids.
63. The polypeptide of claim 5, wherein the amino acid sequence of said truncated neublastin polypeptide comprises amino acids 36-140 of SEQ ID NO:9.
64. The polypeptide of claim 5, wherein the amino acid sequence of said truncated neublastin polypeptide consists essentially of 105 amino acids.
65. The polypeptide of claim 5, wherein the amino acid sequence of said truncated neublastin polypeptide comprises amino acids 35-140 of SEQ ID NO:9.
66. The polypeptide of claim 5, wherein the amino acid sequence of said truncated neublastin polypeptide consists essentially of 106 amino acids.
67. The polypeptide of claim 5, wherein the amino acid sequence of said truncated neublastin polypeptide comprises amino acids 34-140 of SEQ ID NO:9.
68. The polypeptide of claim 5, wherein the amino acid sequence of said truncated neublastin polypeptide consists essentially of 107 amino acids.
69. The polypeptide of claim 5, wherein the amino acid sequence of said truncated neublastin polypeptide comprises amino acids 33-140 of SEQ ID NO:9.

70. The polypeptide of claim 5, wherein the amino acid sequence of said truncated neublastin polypeptide consists essentially of 108 amino acids.
71. The polypeptide of claim 5, wherein the amino acid sequence of said truncated neublastin polypeptide comprises amino acids 32-140 of SEQ ID NO:9.
72. The polypeptide of claim 5, wherein the amino acid sequence of said truncated neublastin polypeptide consists essentially of 109 amino acids.
73. The polypeptide of claim 5, wherein the amino acid sequence of said truncated neublastin polypeptide comprises amino acids 31-140 of SEQ ID NO:9.
74. The polypeptide of claim 5, wherein the amino acid sequence of said truncated neublastin polypeptide consists essentially of 110 amino acids.
75. The polypeptide of claim 5, wherein the amino acid sequence of said truncated neublastin polypeptide comprises amino acids 30-140 of SEQ ID NO:9.
76. The polypeptide of claim 5, wherein the amino acid sequence of said truncated neublastin polypeptide consists essentially of 111 amino acids.
77. The polypeptide of claim 5, wherein the amino acid sequence of said truncated neublastin polypeptide comprises amino acids 29-140 of SEQ ID NO:9.
78. The polypeptide of claim 5, wherein the amino acid sequence of said truncated neublastin polypeptide consists essentially of 112 amino acids.
79. Any one polypeptide of claims 55-78, wherein the amino acid sequence of said truncated neublastin polypeptide is at least 80% homologous to the corresponding sequence in SEQ ID NO:9.

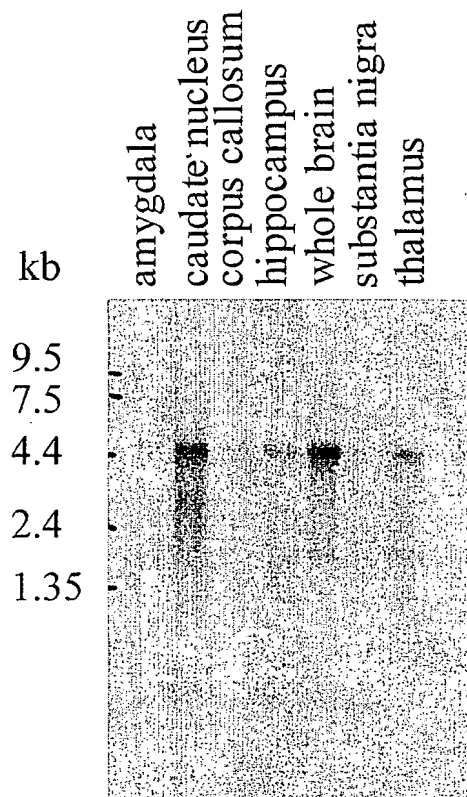


80. The Any one polypeptide of claims 55-78, wherein the amino acid sequence of said neublastin polypeptide is at least 90% homologous to the corresponding sequence in SEQ ID NO:9.
81. Any one polypeptide of claims 55-78, wherein the amino acid sequence of said neublastin polypeptide is at least 95% homologous to the corresponding sequence in SEQ ID NO:9.
82. Any one polypeptide of claims 55-78, wherein the amino acid sequence of said truncated neublastin polypeptide is at least 99% homologous to the corresponding sequence in SEQ ID NO:9.

**A.**



**B.**



Figs. 1A and 1B

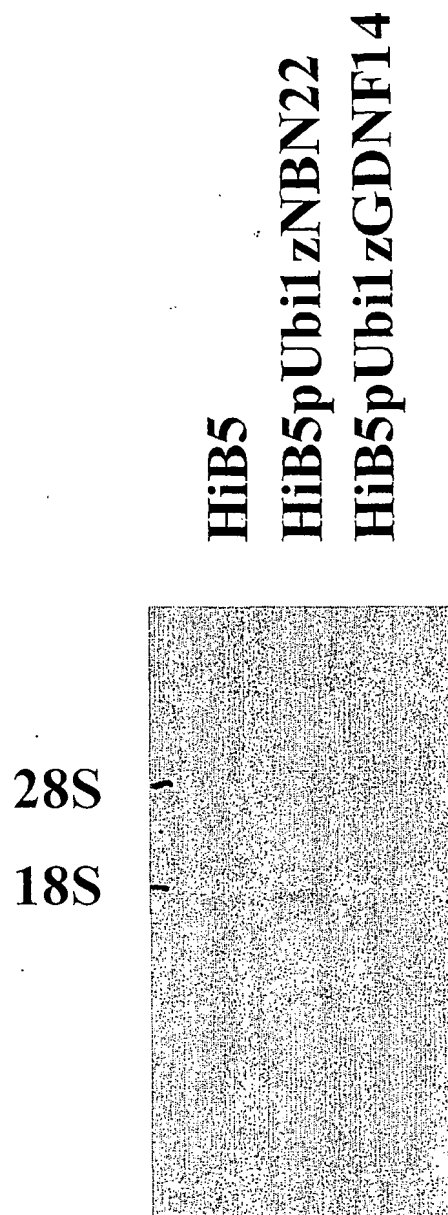


Fig. 2

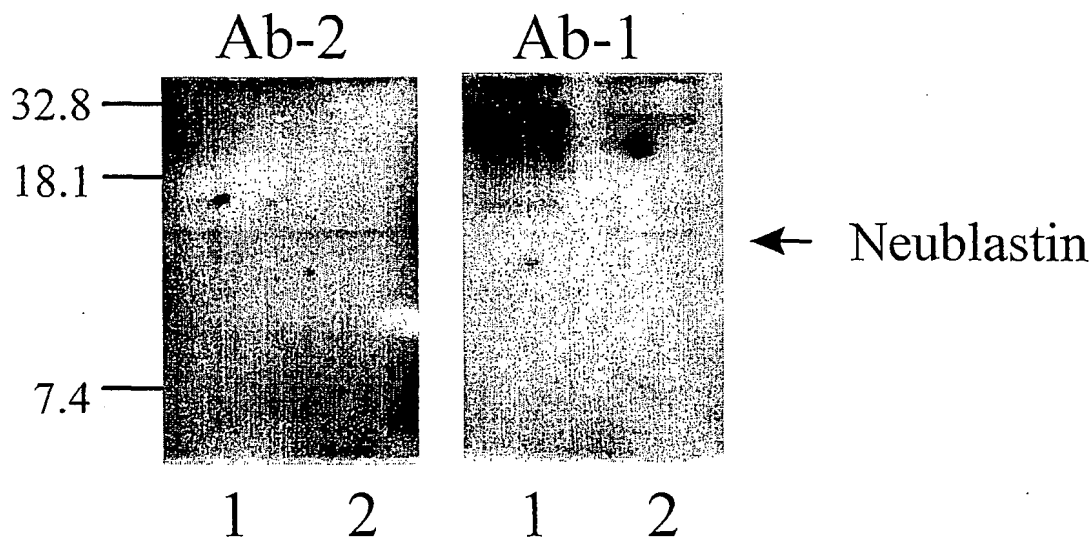
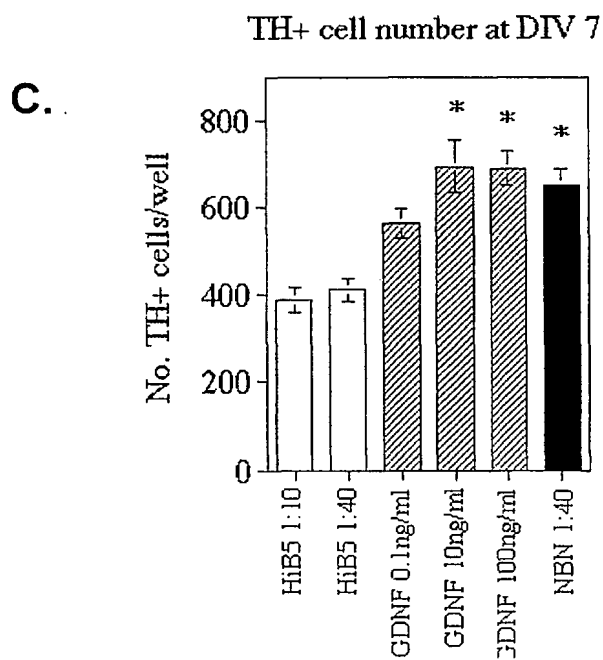
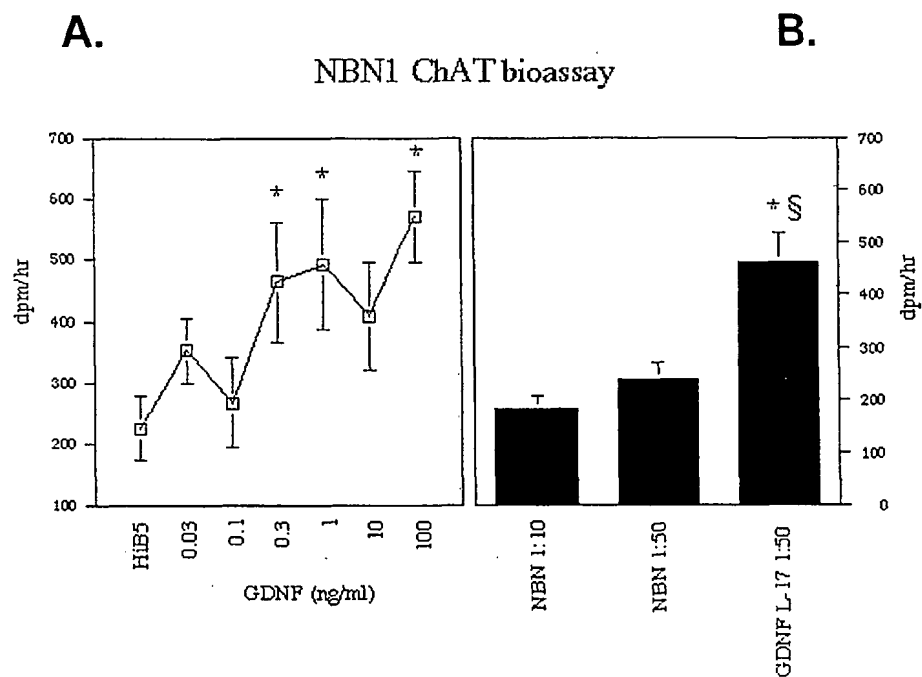
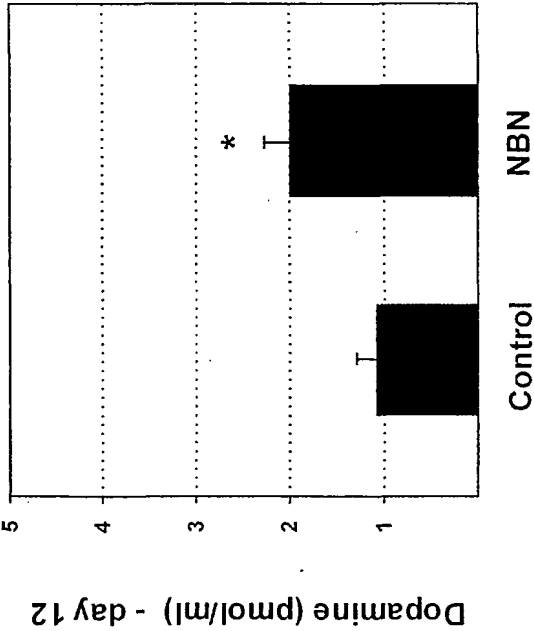


Fig. 3

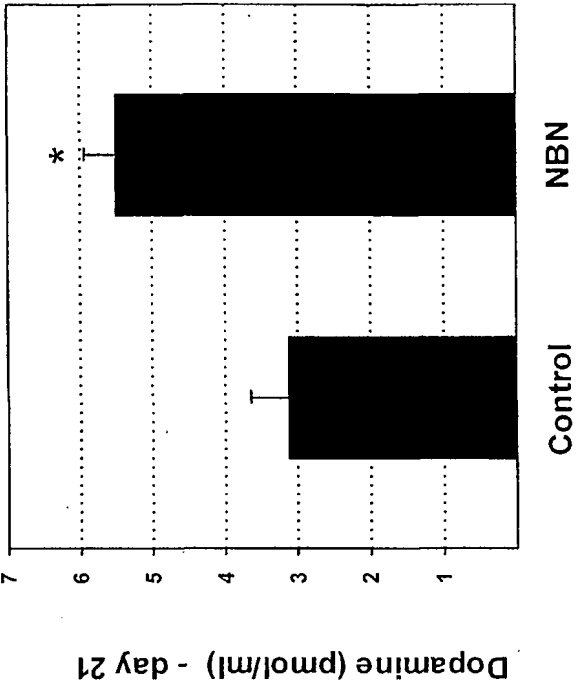


Figs. 4A, 4B and 4C

A.



B.



Figs. 5A and 5B

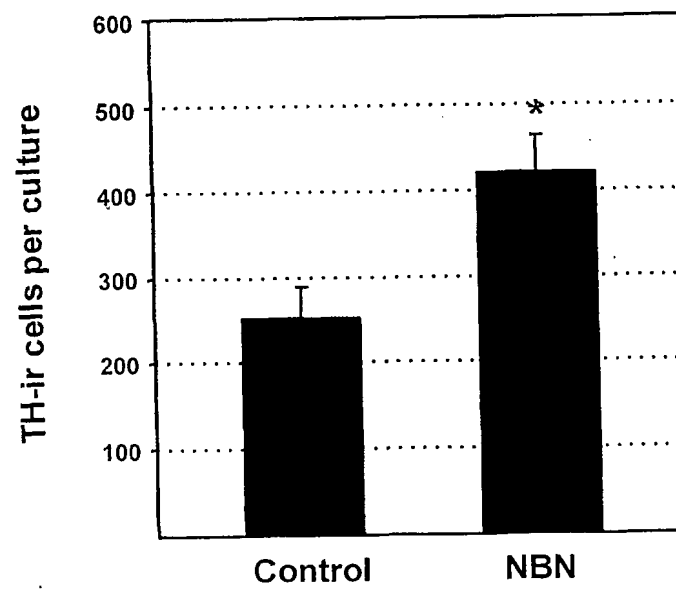
**C.**

Fig. 5C

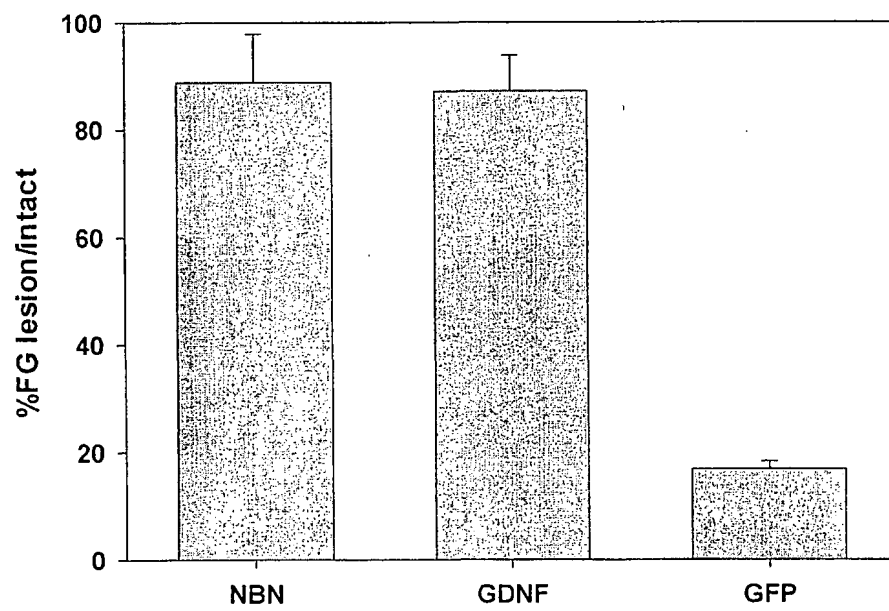
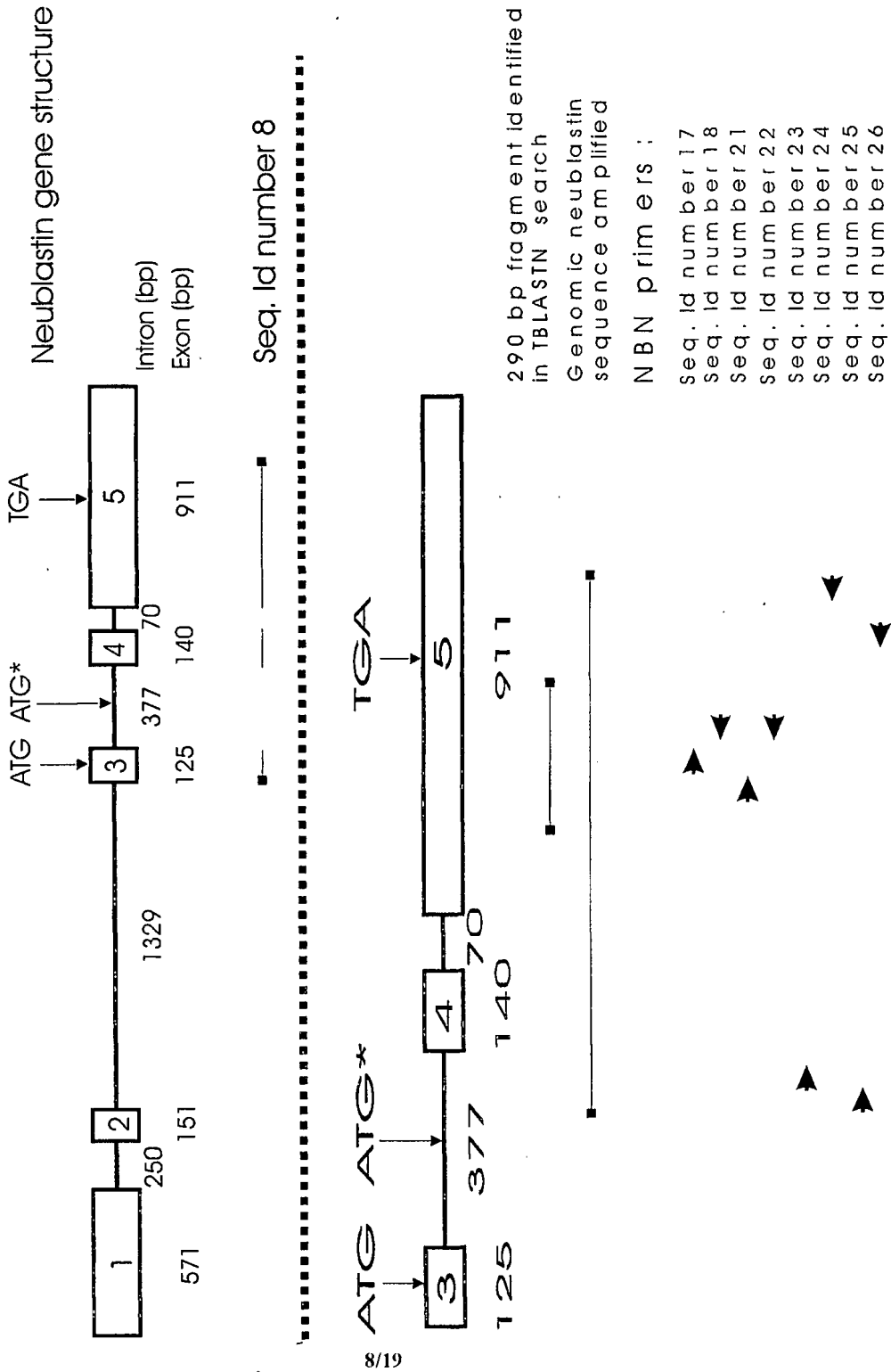
**In vivo effects of NBN on nigral dopamine neurons****Fig. 6**



FIG. 7



Alignment of Neublastin primers used in Rapid-Screen with  
homologous regions in other GDNF ligands

5' -C CTG GCC AGC CTA CTG GG-3'	SEQ ID No 17
G CTG GCC CGG CTG CAG GG	persephin
G CTG CGA CGA CTG CGC CA	neurturin
A TTG AAA AAC TTA TCC AG	GDNF

5' -AA GGA GAC CGC       TTC GTA GCG-3'	SEQ ID No 18
TA GGC CAC GTC       GGT GTA GCG	persephin
AA GGA CAC CTC GTC CTC GTA GGC	neurturin
AA CGA CAG GTC ATC ATC AAA GGC	GDNF

conserved nucleotides shown in bold

**Fig. 8**

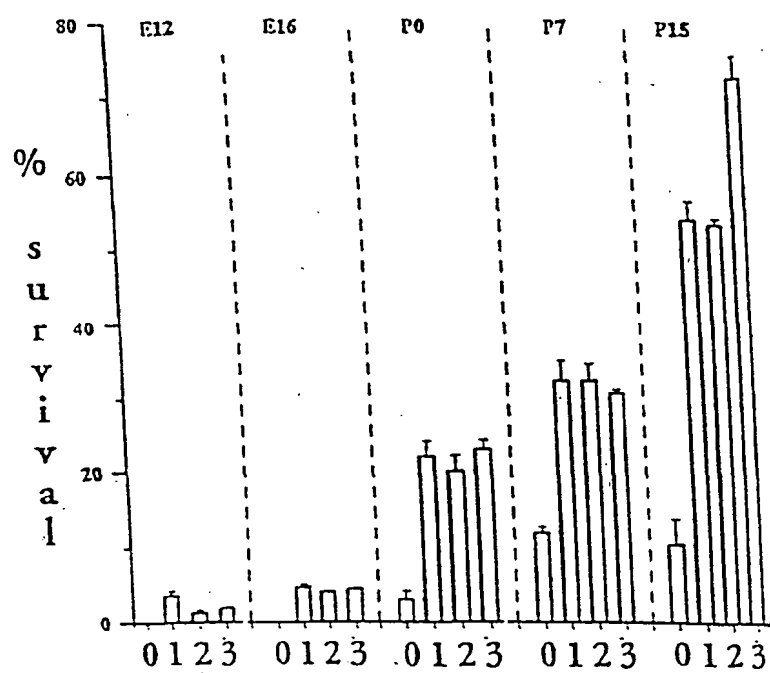


Fig. 9

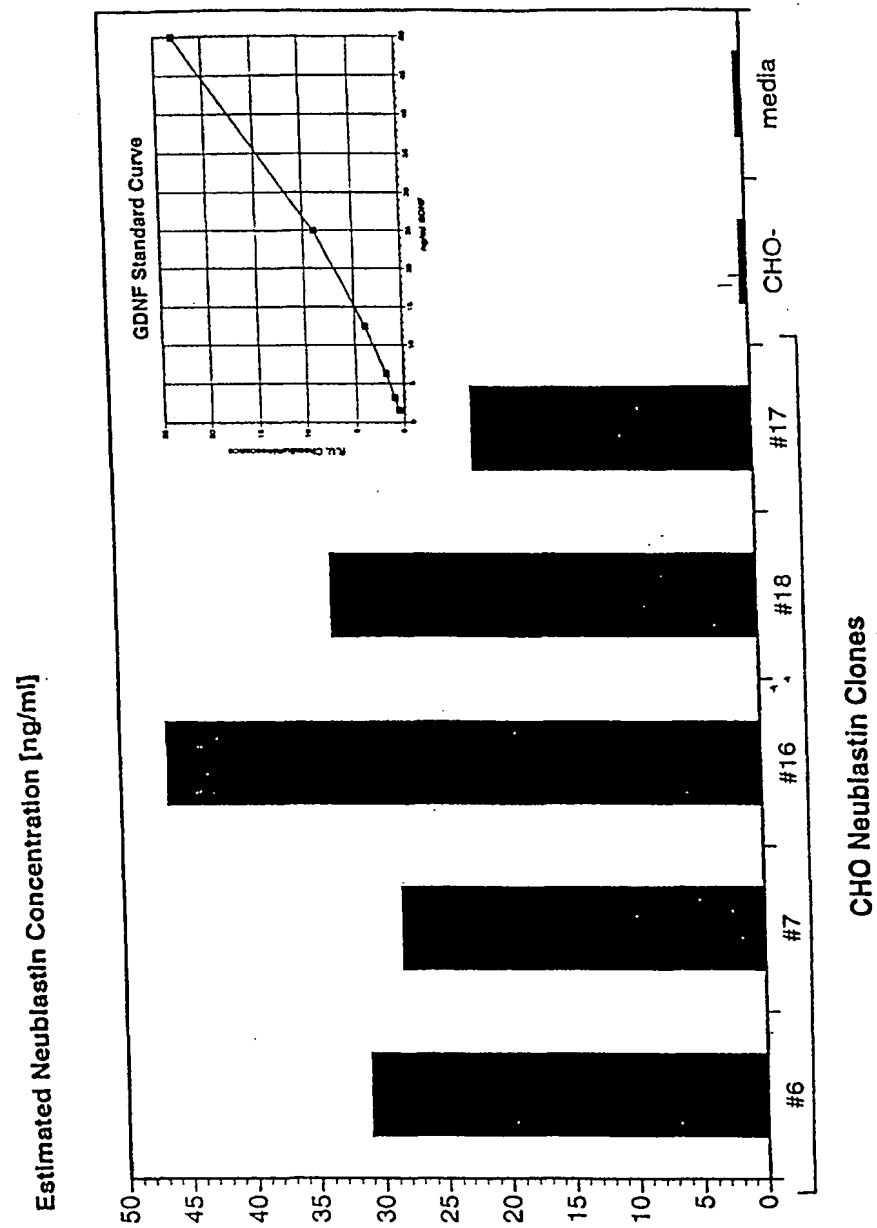


Fig. 10

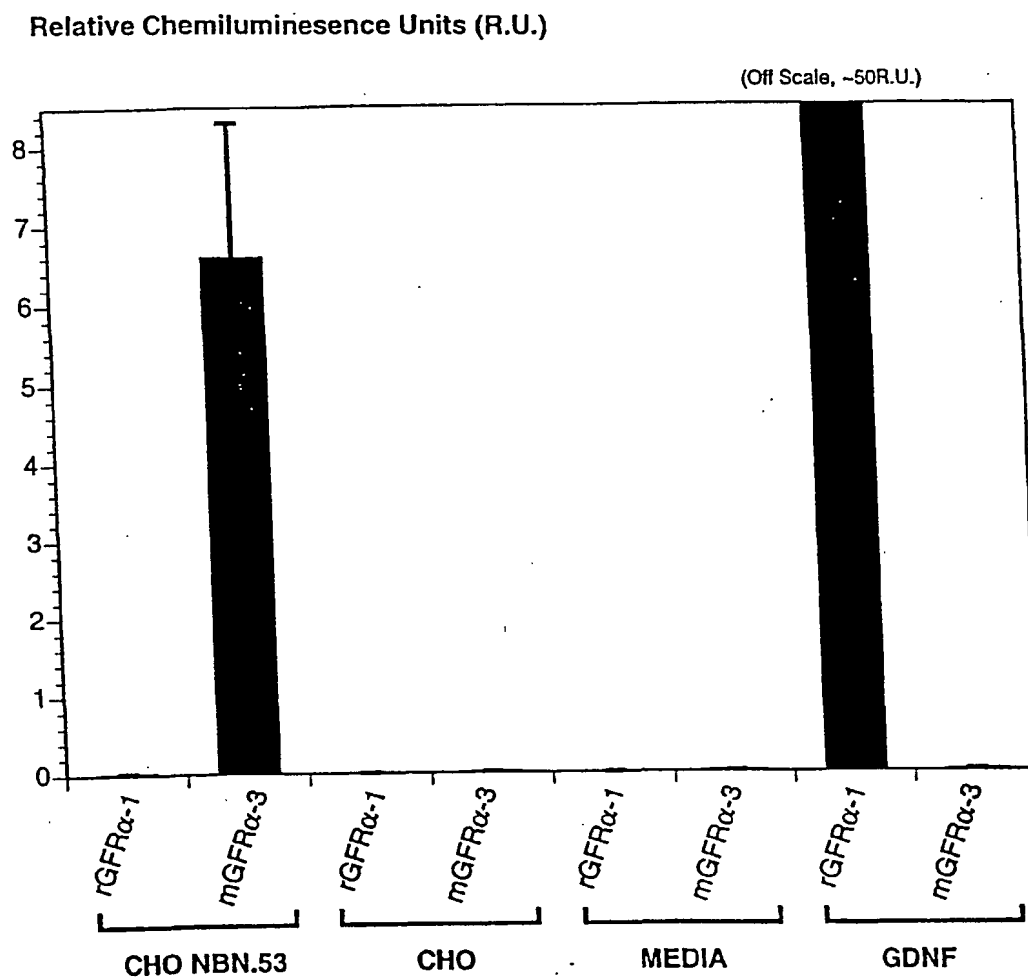
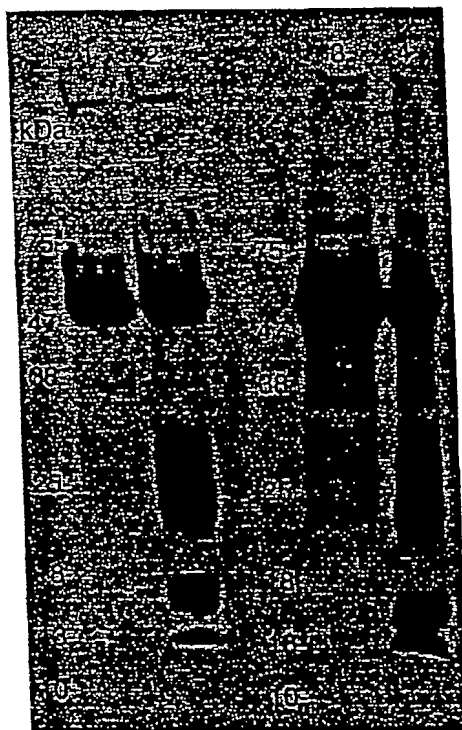
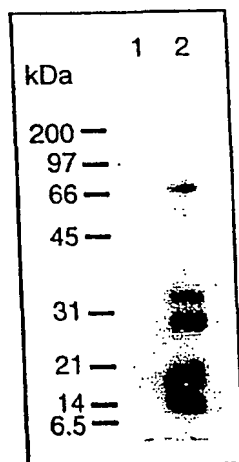


Fig. 11



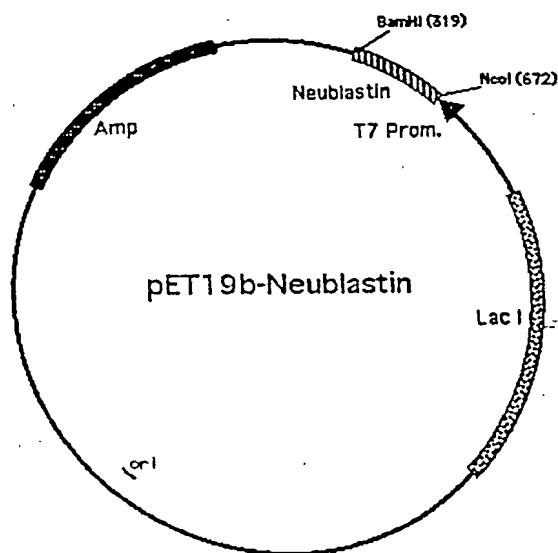
1. Control medium stained with R30 anti-peptide antibody
2. Neublastin containing conditioned medium stained with R30 anti-peptide antibody
3. Control medium stained with R31 anti-peptide antibody
4. Neublastin containing conditioned medium stained with R31 anti-peptide antibody

Fig. 12



Extraction of neublastin by affinity-binding on RETL3-Ig  
Lane 1: bound from CHO control conditioned media  
Lane 2: bound from neublastin overexpressing CHO conditioned media

Fig. 13



## Neublastin Syngene

NcoI (318)

```

316 TACCATGGCT GGAGGACCGG GATCTCGTGC TCGTGCAGCA GGAGCACGTG GCTGTCGTCT
    ATCCTACCGA CCTCCTGGCC CTAGAGCACG AGCACGTCGT CCTCGTGCAC CACACAGAGA
    1 M A G G P G S R A R A A G A R G C R L

376 GCGTTCTCAA CTAGTGCCGG TCGTGCACT CGGACTGGGA CACCGTTCCG ACGAACTAGT
    CGCAAGAGTT GATCACGGCC ACGCACGTGA GCCTGACCCT GTGGCAAGGC TGCTTGATCA
    19 R S Q L V P V R A L G L G H R S D E L V

436 ACGTTTTCGT TTTGTTCAG GATCTTGTCG TCGTGCACGT TCTCCGCATG ATCTATCTCT
    TGCAAAAGCA AAAACAAGTC CTAGAACAGC AGCACGTGCA AGAGGCGTAC TAGATAGAGA
    39 R F R F C S G S C R R A R S P H D L S L

496 AGCATCTCTA CTAGGAGCCG GAGCACTAAG ACCGCCGCCG GGATCTAGAC CTGTATCTCA
    TCGTAGAGAT GATCCTCGGC CTCGTGATTC TGGCGGCGGC CCTAGATCTG GACATAGAGT
    59 A S L L G A G A L R P P P G S R P V S Q

556 ACCTTGTTGT AGACCTACTA GATACGAAGC AGTATCTTTC ATGGACGTAA ACTCTACATG
    TGGAAACAAC TCTGGATGAT CTATGCTTCG TCATAGAAAG TACCTGCATT TGAGATGTAC
    79 P C C R P T R Y E A V S F M D V N S T W

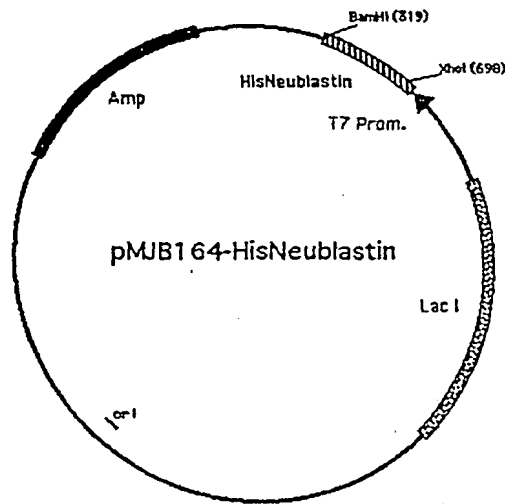
616 GAGAACCGTA GATAGACTAT CTGCAACCGC ATGTGGCTGT CTAGGATGAT AATAGGGATC
    CTCTTGGCAT CTATCTGATA GACGTTGGCG TACACCGACA GATCCTACTA TTATCCCTAG
    99 R T V D R L S A T A C G C L G . . .

676 CGGCT
    GCCGA
  
```

BamHI (671)

Fig. 14





## HisNeubl原因

XhoI (340)

```

301 TACCATGGGC CATCATCATC ATCATCATCA TCATCATCAC TCGAGCGGCC ATATCGACGA
    ATCTTACCCG GTAGTAGTAG TAGTAGTAGT AGTAGTAGTG AGCTCGCCGG TATAGCTGCT
    1▶ M G H H H H H H H H H S S G H I D D

361 CGACGACAAG GCTGGAGGAC CGGGATCTCG TGCTCGTGCA GCAGGAGCAC GTGGCTGTCTG
    SCTGCTGTTC CGACCTCCTG GCCCTAGAGC ACGAGCACGT CGTCCTCGTG CACCGACAGC
    19▶ D D K A G G P G S R A R A A G A R G C R

421 TCTGCGTTCT CAACTAGTGC CGGTGCGTGC ACTCGGACTG GGACACCGTT CCGACGAACT
    AGACGCAAGA GTTGATCACG GCCACGCACG TGAGCCTGAC CCTGTGGCAA GGCTGCTTGA
    39▶ L R S Q L V P V R A L G L G H R S D E L

481 AGTACGTTTT CGTTTTTGTG CAGGATCTTG TCGTCGTGCA CGTTCTCCGC ATGATCTATC
    TCATGCAAAA GCAAAAACAA GTCCTAGAAC AGCAGCACGT GCAAGAGGCG TACTAGATAG
    59▶ V R F R F C S G S C R R A R S P H D L S

541 TCTAGCATCT CTAGTAGGAG CCGGAGCACT AAGACCGCCG CCGGGATCTA GACCTGTATC
    AGATCGTAGA GATGATCCTC GGCCTCGTGA TTCTGGCGGC GGCCCTAGAT CTGGACATAG
    79▶ L A S L L G A G A L R P P P G S R P V S

601 TCAACCTTGT TGTAGACCTA CTAGATACGA AGCAGTATCT TTCATGGACG TAAACTCTAC
    AGTTGGAACA ACATCTGGAT GATCTATGCT TCGTCATAGA AAGTACCTGC ATTTGAGATG
    99▶ Q P C C R P T R Y E A V S F M D V N S T

661 ATGGAGAACC GTAGATAGAC TATCTGCAAC CGCATGTGGC TGTCTAGGAT GATAATAGGG
    TACCTCTTGG CATCTATCTG ATAGACGTTG GCGTACACCG ACAGATCCTA CTATTATCCC
    119▶ W R T V D R L S A T A C G C L G . .

721 ATCCGGCTGC TAACAAAGCC CG
    TAGGCCGACG ATTGTTTCGG GC
  
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BamHI (719)

Fig. 15

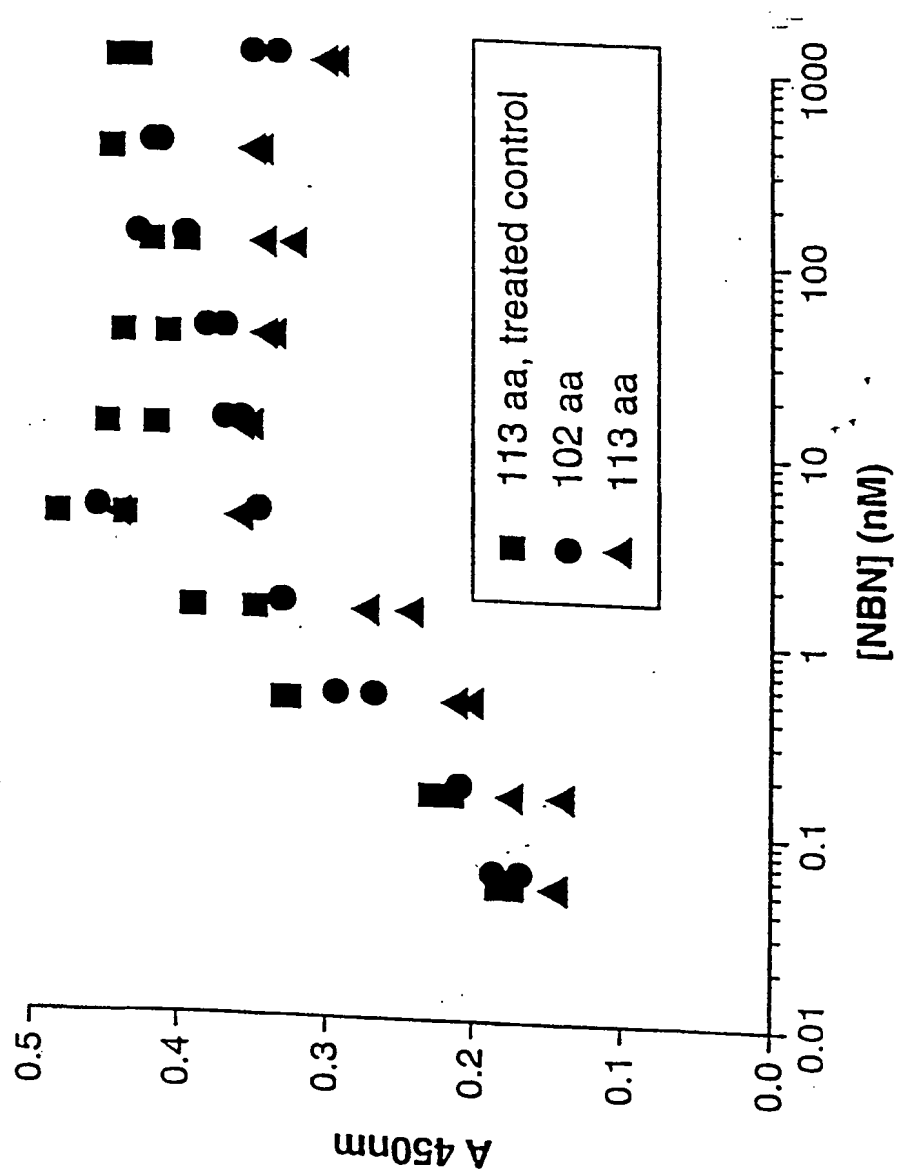


Fig. 16

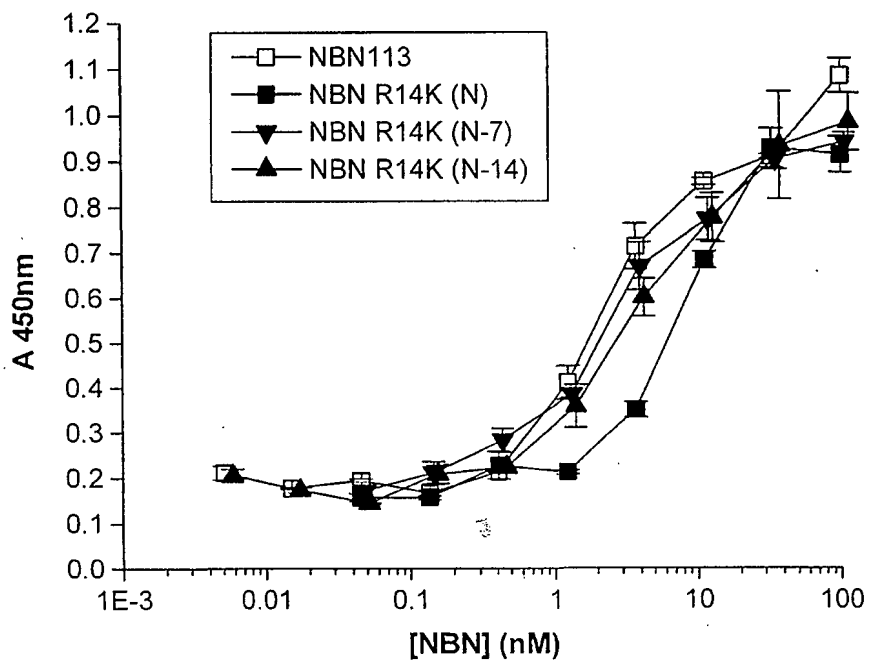


FIG. 17

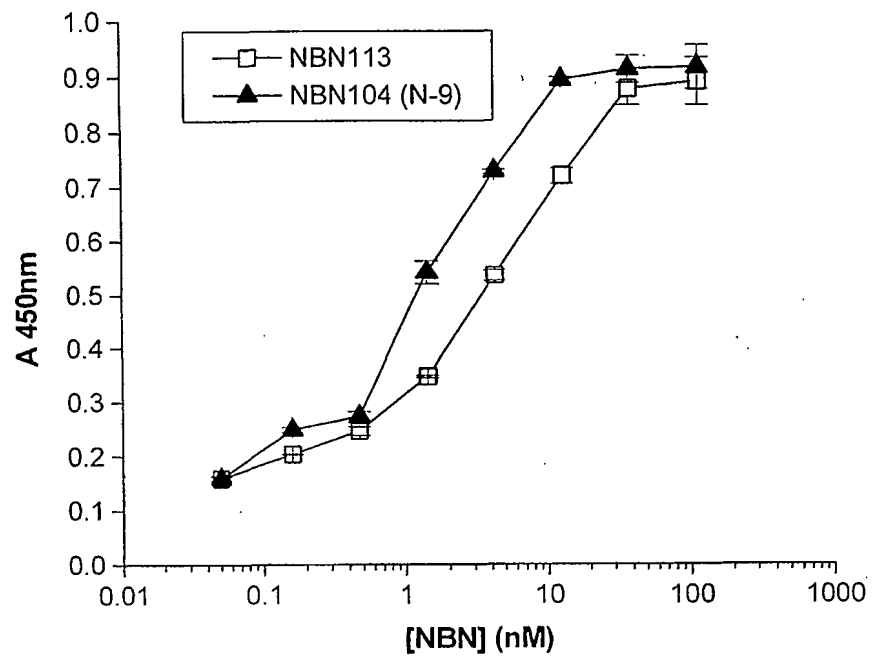


FIG. 18

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Biogen, Inc.  
Johansen, Teit E.  
Wen-Yee Sah, Dinah

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tccccaagcc cacctgggtg cccctcttct cctgaggct ccacttggtc tctccgcgc 119

atg cct gcc ctg tgg ccc acc ctg gcc gct ctg gct ctg ctg agc agc 167  
Met Pro Ala Leu Trp Pro Thr Leu Ala Ala Leu Ala Leu Leu Ser Ser  
-95 -90 -85 -80

gtc gca gag gcc tcc ctg ggc tcc gcg ccc cgc agc cct gcc ccc cgc 215  
Val Ala Glu Ala Ser Leu Gly Ser Ala Pro Arg Ser Pro Ala Pro Arg  
-75 -70 -65

gaa ggc ccc ccg cct gtc ctg gcg tcc ccc gcc ggc cac ctg ccg ggg 263  
Glu Gly Pro Pro Pro Val Leu Ala Ser Pro Ala Gly His Leu Pro Gly  
-60 -55 -50

gga cgc acg gcc cgc tgg tgc agt gga aga gcc cgg cgg ccg cgc cgc 311  
Gly Arg Thr Ala Arg Trp Cys Ser Gly Arg Ala Arg Arg Pro Arg Arg  
-45 -40 -35

aga cac ttc tcg gcc cgc gcc ccc gcc gcc tgc acc ccc atc tgc tct 359  
Arg His Phe Ser Ala Arg Ala Pro Ala Ala Cys Thr Pro Ile Cys Ser  
-30 -25 -20

tcc ccg cgg gtc cgc gcg gcg cgg ctg ggg ggc cgg gca gcg cgc tcg 407  
Ser Pro Arg Val Arg Ala Ala Arg Leu Gly Gly Arg Ala Ala Arg Ser  
-15 -10 -5 -1 1

ggc agc ggg ggc gcg ggg tgc cgc ctg cgc tcg cag ctg gtg ccg gtg 455  
Gly Ser Gly Gly Ala Gly Cys Arg Leu Arg Ser Gln Leu Val Pro Val  
5 10 15

cgc gcg ctc ggc ctg ggc cac cgc tcc gac gag ctg gtg cgt ttc cgc 503  
Arg Ala Leu Gly Leu Gly His Arg Ser Asp Glu Leu Val Arg Phe Arg  
20 25 30

ttc tgc acc ggc tcc tgc ccg cgc gcg cgc tct cca cac gac ctc agc 551

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Phe Cys Thr Gly Ser Cys Pro Arg Ala Arg Ser Pro His Asp Leu Ser
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ctg gcc agc cta ctg ggc gcc ggg gcc ctg cga ccg ccc ccg ggc tcc 599
Leu Ala Ser Leu Leu Gly Ala Gly Ala Leu Arg Pro Pro Pro Gly Ser
   50              55              60              65

cgg ccc gtc agc cag ccc tgc tgc cga ccc acg cgc tac gaa gcg gtc 647
Arg Pro Val Ser Gln Pro Cys Cys Arg Pro Thr Arg Tyr Glu Ala Val
              70              75              80

tcc ttc atg gac gtc aac agc acc tgg aga acc gtg gac cgc ctc tcc 695
Ser Phe Met Asp Val Asn Ser Thr Trp Arg Thr Val Asp Arg Leu Ser
              85              90              95

gcc acc gcc tgc ggc tgc ctg ggc tgagggctcg ctccagggct ttgcagactg 749
Ala Thr Ala Cys Gly Cys Leu Gly
   100              105

gacccttacc ggtggtctctt cctgcctggg accctcccgc agagtccac tagccagcgg 809

cctcagccag ggacgaaggc ctcaaagctg agaggcccct gccggtgggt gatgga 865

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Glu Gly Pro Pro Pro Val Leu Ala Ser Pro Ala Gly His Leu Pro Gly
              -60              -55              -50

Gly Arg Thr Ala Arg Trp Cys Ser Gly Arg Ala Arg Arg Pro Arg Arg
              -45              -40              -35

Arg His Phe Ser Ala Arg Ala Pro Ala Ala Cys Thr Pro Ile Cys Ser
              -30              -25              -20

Ser Pro Arg Val Arg Ala Ala Arg Leu Gly Gly Arg Ala Ala Arg Ser
-15              -10              -5              -1 1

Gly Ser Gly Gly Ala Gly Cys Arg Leu Arg Ser Gln Leu Val Pro Val
              5              10              15

Arg Ala Leu Gly Leu Gly His Arg Ser Asp Glu Leu Val Arg Phe Arg
              20              25              30

Phe Cys Thr Gly Ser Cys Pro Arg Ala Arg Ser Pro His Asp Leu Ser
   35              40              45

Leu Ala Ser Leu Leu Gly Ala Gly Ala Leu Arg Pro Pro Pro Gly Ser
   50              55              60              65

Arg Pro Val Ser Gln Pro Cys Cys Arg Pro Thr Arg Tyr Glu Ala Val
              70              75              80

Ser Phe Met Asp Val Asn Ser Thr Trp Arg Thr Val Asp Arg Leu Ser
              85              90              95

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 <223> DISULFID: Cys107-Cys107 interchain disulfide  
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    Met Pro Gly Leu Ile Ser Ala Arg Gly Gln Pro Leu Leu Glu
          -95          -90          -85

gtc ctt cct ccc caa gcc cac ctg ggt gcc ctc ttt ctc cct gag gct      96
Val Leu Pro Pro Gln Ala His Leu Gly Ala Leu Phe Leu Pro Glu Ala
          -80          -75          -70

cca ctt ggt ctc tcc gcg cag cct gcc ctg tgg ccc acc ctg gcc gct      144
Pro Leu Gly Leu Ser Ala Gln Pro Ala Leu Trp Pro Thr Leu Ala Ala
          -65          -60          -55

ctg gct ctg ctg agc agc gtc gca gag gcc tcc ctg ggc tcc gcg ccc      192
Leu Ala Leu Leu Ser Ser Val Ala Glu Ala Ser Leu Gly Ser Ala Pro
          -50          -45          -40

cgc agc cct gcc ccc cgc gaa ggc ccc ccg cct gtc ctg gcg tcc ccc      240
Arg Ser Pro Ala Pro Arg Glu Gly Pro Pro Pro Val Leu Ala Ser Pro
          -35          -30          -25

gcc ggc cac ctg ccg ggg gga cgc acg gcc cgc tgg tgc agt gga aga      288
Ala Gly His Leu Pro Gly Gly Arg Thr Ala Arg Trp Cys Ser Gly Arg
          -15          -10          -5

gcc cgg ccg ccg ccg ccg cag cct tct cgg ccc gcg ccc ccg ccg cct      336
Ala Arg Arg Pro Pro Pro Gln Pro Ser Arg Pro Ala Pro Pro Pro Pro
          -1    1          5          10

gca ccc cca tct gct ctt ccc cgc ggg ggc cgc gcg gcg ccg gct ggg      384
Ala Pro Pro Ser Ala Leu Pro Arg Gly Gly Arg Ala Ala Arg Ala Gly
          15          20          25

ggc ccg ggc aac cgc gct ccg gca gcg ggg gcg ccg ggc tgc cgc ctg      432
Gly Pro Gly Asn Arg Ala Arg Ala Ala Gly Ala Arg Gly Cys Arg Leu
          30          35          40          45

cgc tgc cag ctg gtg ccg gtg cgc gcg ctc ggc ctg ggc cac cgc tcc      480
Arg Ser Gln Leu Val Pro Val Arg Ala Leu Gly Leu Gly His Arg Ser
          50          55          60

gac gag ctg gtg cgt ttc cgc ttc tgc agc ggc tcc tgc cgc cgc gcg      528
Asp Glu Leu Val Arg Phe Arg Phe Cys Ser Gly Ser Cys Arg Arg Ala
          65          70          75

cgc tct cca cac gac ctc agc ctg gcc agc cta ctg ggc gcc ggg gcc      576
Arg Ser Pro His Asp Leu Ser Leu Ala Ser Leu Leu Gly Ala Gly Ala
          80          85          90

ctg cga ccg ccc ccg ggc tcc ccg ccc gtc agc cag ccc tgc tgc cga      624
Leu Arg Pro Pro Pro Gly Ser Arg Pro Val Ser Gln Pro Cys Cys Arg
          95          100          105

ccc acg cgc tac gaa gcg gtc tcc ttc atg gac gtc aac agc acc tgg      672
Pro Thr Arg Tyr Glu Ala Val Ser Phe Met Asp Val Asn Ser Thr Trp
          110          115          120          125

aga acc gtg gac cgc ctc tcc gcc aac ccc tgc ggc tgc ctg ggc      717
Arg Thr Val Asp Arg Leu Ser Ala Asn Pro Cys Gly Cys Leu Gly
          130          135          140

tgagggctcg ctccagggct ttgcagactg gacccttacc ggtggctctt cctgectggg 777

accctccgcg agagtccac tagccagcgg cctcagccag ggacgaaggc ctcaaagctg 837

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861

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&lt;213&gt; Homo sapiens

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Pro Pro Gln Ala His Leu Gly Ala Leu Phe Leu Pro Glu Ala Pro Leu
      -80                      -75                      -70

Gly Leu Ser Ala Gln Pro Ala Leu Trp Pro Thr Leu Ala Ala Leu Ala
-65                      -60                      -55                      -50

Leu Leu Ser Ser Val Ala Glu Ala Ser Leu Gly Ser Ala Pro Arg Ser
                      -45                      -40                      -35

Pro Ala Pro Arg Glu Gly Pro Pro Pro Val Leu Ala Ser Pro Ala Gly
                      -30                      -25                      -20

His Leu Pro Gly Gly Arg Thr Ala Arg Trp Cys Ser Gly Arg Ala Arg
      -15                      -10                      -5

Arg Pro Pro Pro Gln Pro Ser Arg Pro Ala Pro Pro Pro Pro Ala Pro
-1   1                      5                      10                      15

Pro Ser Ala Leu Pro Arg Gly Gly Arg Ala Ala Arg Ala Gly Gly Pro
      20                      25                      30

Gly Asn Arg Ala Arg Ala Ala Gly Ala Arg Gly Cys Arg Leu Arg Ser
      35                      40                      45

Gln Leu Val Pro Val Arg Ala Leu Gly Leu Gly His Arg Ser Asp Glu
      50                      55                      60

Leu Val Arg Phe Arg Phe Cys Ser Gly Ser Cys Arg Arg Ala Arg Ser
      65                      70                      75

Pro His Asp Leu Ser Leu Ala Ser Leu Leu Gly Ala Gly Ala Leu Arg
      80                      85                      90                      95

Pro Pro Pro Gly Ser Arg Pro Val Ser Gln Pro Cys Cys Arg Pro Thr
      100                      105                      110

Arg Tyr Glu Ala Val Ser Phe Met Asp Val Asn Ser Thr Trp Arg Thr
      115                      120                      125

Val Asp Arg Leu Ser Ala Asn Pro Cys Gly Cys Leu Gly
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&lt;211&gt; 140

&lt;212&gt; PRT

&lt;213&gt; Homo sapiens

&lt;220&gt;

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and Xaa at position 135 designates Ala or Pro

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     1                    5                    10                    15  
 Ser Ala Leu Pro Arg Gly Gly Arg Ala Ala Arg Ala Gly Gly Pro Gly  
           20                    25                    30  
 Asn Arg Ala Arg Ala Ala Gly Ala Arg Gly Cys Arg Leu Arg Ser Gln  
           35                    40                    45  
 Leu Val Pro Val Arg Ala Leu Gly Leu Gly His Arg Ser Asp Glu Leu  
           50                    55                    60  
 Val Arg Phe Arg Phe Cys Ser Gly Ser Cys Arg Arg Ala Arg Ser Pro  
           65                    70                    75                    80  
 His Asp Leu Ser Leu Ala Ser Leu Leu Gly Ala Gly Ala Leu Arg Pro  
                     85                    90                    95  
 Pro Pro Gly Ser Arg Pro Val Ser Gln Pro Cys Cys Arg Pro Thr Arg  
                     100                    105                    110  
 Tyr Glu Ala Val Ser Phe Met Asp Val Asn Ser Thr Trp Arg Thr Val  
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           20                    25                    30  
 Leu Gly His Arg Ser Asp Glu Leu Val Arg Phe Arg Phe Cys Ser Gly  
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 Gln Pro Cys Cys Arg Pro Thr Arg Tyr Glu Ala Val Ser Phe Met Asp  
                     85                    90                    95  
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 Gly Cys Leu Gly  
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<210> 7

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 and Xaa at position 108 designates Ala or Pro  
  
 <400> 7  
 Ala Gly Gly Pro Gly Asn Arg Ala Arg Ala Ala Gly Ala Arg Gly Cys  
 1 5 10 15  
 Arg Leu Arg Ser Gln Leu Val Pro Val Arg Ala Leu Gly Leu Gly His  
 20 25 30  
 Arg Ser Asp Glu Leu Val Arg Phe Arg Phe Cys Ser Gly Ser Cys Arg  
 35 40 45  
 Arg Ala Arg Ser Pro His Asp Leu Ser Leu Ala Ser Leu Leu Gly Ala  
 50 55 60  
 Gly Ala Leu Arg Pro Pro Pro Gly Ser Arg Pro Val Ser Gln Pro Cys  
 65 70 75 80  
 Cys Arg Pro Thr Arg Tyr Glu Ala Val Ser Phe Met Asp Val Asn Ser  
 85 90 95  
 Thr Trp Arg Thr Val Asp Arg Leu Ser Ala Xaa Xaa Cys Gly Cys Leu  
 100 105 110

Gly

<210> 8  
 <211> 861  
 <212> DNA  
 <213> Homo sapiens

<220>  
 <221> CDS  
 <222> (58) .. (717)

<220>  
 <221> 5'UTR  
 <222> (1) .. (57)

<220>  
 <221> 3'UTR  
 <222> (718) .. (861)

<220>  
 <221> sig\_peptide  
 <222> (58) .. (174)

<220>  
 <221> mat\_peptide  
 <222> (298) .. (717)

<220>  
 <221> mat\_peptide  
 <222> (370) .. (717)

<220>  
 <221> mat\_peptide

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<222> (379)..(717)

<220>
<221> misc_structure
<222> (661)..(663)
<223> CARBOHYD: glycosylated asparagine at Asn122

<220>
<221> misc_structure
<222> (424)..(621)
<223> DISULFID: Gly43-Gly108 disulfide bridge

<220>
<221> misc_structure
<222> (505)..(705)
<223> DISULFID: Gly70-Gly136 disulfide bridge

<220>
<221> misc_structure
<222> (517)..(711)
<223> DISULFID: Gly74-Gly138 disulfide bridge

<220>
<221> misc_structure
<222> (616)..(618)
<223> DISULFID: Gly107-Gly107 interchain disulfide
      bridge

<400> 8
aggagggtgg gggaacagct caacaatggc tgatgggcgc tcctggtgtt gatagag      57

atg gaa ctt gga ctt gga ggc ctc tcc acg ctg tcc cac tgc ccc tgg      105
Met Glu Leu Gly Leu Gly Gly Leu Ser Thr Leu Ser His Cys Pro Trp
-80              -75              -70              -65

cct agg cgg cag cct gcc ctg tgg ccc acc ctg gcc gct ctg gct ctg      153
Pro Arg Arg Gln Pro Ala Leu Trp Pro Thr Leu Ala Ala Leu Ala Leu
-60              -55              -50

ctg agc agc gtc gca gag gcc tcc ctg ggc tcc gcg ccc cgc agc cct      201
Leu Ser Ser Val Ala Glu Ala Ser Leu Gly Ser Ala Pro Arg Ser Pro
-45              -40              -35

gcc ccc cgc gaa ggc ccc ccg cct gtc ctg gcg tcc ccc gcc ggc cac      249
Ala Pro Arg Glu Gly Pro Pro Pro Val Leu Ala Ser Pro Ala Gly His
-30              -25              -20

ctg ccg ggg gga cgc acg gcc cgc tgg tgc agt gga aga gcc cgg cgg      297
Leu Pro Gly Gly Arg Thr Ala Arg Trp Cys Ser Gly Arg Ala Arg Arg
-15              -10              -5              -1

ccg ccg ccg cag cct tct ccg ccc gcg ccc ccg ccg cct gca ccc cca      345
Pro Pro Pro Gln Pro Ser Arg Pro Ala Pro Pro Pro Pro Ala Pro Pro
1              5              10              15

tct gct ctt ccc cgc ggg ggc cgc gcg gcg cgg gct ggg ggc ccg ggc      393
Ser Ala Leu Pro Arg Gly Gly Arg Ala Ala Arg Ala Gly Gly Pro Gly
20              25              30

agc cgc gct cgg gca gcg ggg gcg cgg ggc tgc cgc ctg cgc tcg cag      441
Ser Arg Ala Arg Ala Ala Gly Ala Arg Gly Cys Arg Leu Arg Ser Gln
35              40              45

ctg gtg ccg gtg cgc gcg ctc ggc ctg ggc cac cgc tcc gac gag ctg      489
Leu Val Pro Val Arg Ala Leu Gly Leu Gly His Arg Ser Asp Glu Leu

```

50                      55                      60  
 gtg cgt ttc cgc ttc tgc agc ggc tcc tgc cgc cgc gcg cgc tct cca 537  
 Val Arg Phe Arg Phe Cys Ser Gly Ser Cys Arg Arg Ala Arg Ser Pro  
 65                      70                      75                      80  
 cac gac ctc agc ctg gcc agc cta ctg ggc gcc ggc gcc ctg cga ccg 585  
 His Asp Leu Ser Leu Ala Ser Leu Leu Gly Ala Gly Ala Leu Arg Pro  
 85                      90                      95  
 ccc ccg ggc tcc cgg ccc gtc agc cag ccc tgc tgc cga ccc acg cgc 633  
 Pro Pro Gly Ser Arg Pro Val Ser Gln Pro Cys Cys Arg Pro Thr Arg  
 100                      105                      110  
 tac gaa gcg gtc tcc ttc atg gac gtc aac agc acc tgg aga acc gtg 681  
 Tyr Glu Ala Val Ser Phe Met Asp Val Asn Ser Thr Trp Arg Thr Val  
 115                      120                      125  
 gac cgc ctc tcc gcc acc gcc tgc ggc tgc ctg ggc tgagggctcg 727  
 Asp Arg Leu Ser Ala Thr Ala Cys Gly Cys Leu Gly  
 130                      135                      140  
 ctccagggct ttgcagactg gacccttacc ggtggctctt cctgcctggg accctccgc 787  
 agagtcccac tagccagcgg cctcagccag ggacgaaggc ctcaaagctg agaggcccct 847  
 accggtgggt gatg 861  
  
 <210> 9  
 <211> 220  
 <212> PRT  
 <213> Homo sapiens  
  
 <400> 9  
 Met Glu Leu Gly Leu Gly Gly Leu Ser Thr Leu Ser His Cys Pro Trp  
 -80                      -75                      -70                      -65  
 Pro Arg Arg Gln Pro Ala Leu Trp Pro Thr Leu Ala Ala Leu Ala Leu  
 -60                      -55                      -50  
 Leu Ser Ser Val Ala Glu Ala Ser Leu Gly Ser Ala Pro Arg Ser Pro  
 -45                      -40                      -35  
 Ala Pro Arg Glu Gly Pro Pro Pro Val Leu Ala Ser Pro Ala Gly His  
 -30                      -25                      -20  
 Leu Pro Gly Gly Arg Thr Ala Arg Trp Cys Ser Gly Arg Ala Arg Arg  
 -15                      -10                      -5                      -1  
 Pro Pro Pro Gln Pro Ser Arg Pro Ala Pro Pro Pro Pro Ala Pro Pro  
 1                      5                      10                      15  
 Ser Ala Leu Pro Arg Gly Gly Arg Ala Ala Arg Ala Gly Gly Pro Gly  
 20                      25                      30  
 Ser Arg Ala Arg Ala Ala Gly Ala Arg Gly Cys Arg Leu Arg Ser Gln  
 35                      40                      45  
 Leu Val Pro Val Arg Ala Leu Gly Leu Gly His Arg Ser Asp Glu Leu  
 50                      55                      60  
 Val Arg Phe Arg Phe Cys Ser Gly Ser Cys Arg Arg Ala Arg Ser Pro  
 65                      70                      75                      80

<400> 11  
Ala Ala Arg Ala Gly Gly Pro Gly Ser Arg Ala Arg Ala Ala Gly Ala

1                      5                      10                      15  
 Arg Gly Cys Arg Leu Arg Ser Gln Leu Val Pro Val Arg Ala Leu Gly  
                             20                      25                      30  
 Leu Gly His Arg Ser Asp Glu Leu Val Arg Phe Arg Phe Cys Ser Gly  
                             35                      40                      45  
 Ser Cys Arg Arg Ala Arg Ser Pro His Asp Leu Ser Leu Ala Ser Leu  
                             50                      55                      60  
 Leu Gly Ala Gly Ala Leu Arg Pro Pro Pro Gly Ser Arg Pro Val Ser  
                             65                      70                      75                      80  
 Gln Pro Cys Cys Arg Pro Thr Arg Tyr Glu Ala Val Ser Phe Met Asp  
                             85                      90                      95  
 Val Asn Ser Thr Trp Arg Thr Val Asp Arg Leu Ser Ala Thr Ala Cys  
                             100                      105                      110  
 Gly Cys Leu Gly  
                             115

<210> 12  
 <211> 113  
 <212> PRT  
 <213> Homo sapiens

<220>  
 <221> CARBOHYD  
 <222> (95)  
 <223> glycosylated asparagine

<400> 12  
 Ala Gly Gly Pro Gly Ser Arg Ala Arg Ala Ala Gly Ala Arg Gly Cys  
                             1                      5                      10                      15  
 Arg Leu Arg Ser Gln Leu Val Pro Val Arg Ala Leu Gly Leu Gly His  
                             20                      25                      30  
 Arg Ser Asp Glu Leu Val Arg Phe Arg Phe Cys Ser Gly Ser Cys Arg  
                             35                      40                      45  
 Arg Ala Arg Ser Pro His Asp Leu Ser Leu Ala Ser Leu Leu Gly Ala  
                             50                      55                      60  
 Gly Ala Leu Arg Pro Pro Pro Gly Ser Arg Pro Val Ser Gln Pro Cys  
                             65                      70                      75                      80  
 Cys Arg Pro Thr Arg Tyr Glu Ala Val Ser Phe Met Asp Val Asn Ser  
                             85                      90                      95  
 Thr Trp Arg Thr Val Asp Arg Leu Ser Ala Thr Ala Cys Gly Cys Leu  
                             100                      105                      110  
 Gly

<210> 13  
 <211> 102  
 <212> DNA  
 <213> Homo sapiens



13

15	20	25	
gtt cta gcc ctg ctg agc tgc gtc aca gaa gct tcc ctg gac cca atg			1106
Val Leu Ala Leu Leu Ser Cys Val Thr Glu Ala Ser Leu Asp Pro Met			
30	35	40	
tcc cgc agc ccc gcc gct cgc gac ggt ccc tca ccg gtc ttg gcg ccc			1154
Ser Arg Ser Pro Ala Ala Arg Asp Gly Pro Ser Pro Val Leu Ala Pro			
45	50	55	60
ccc acg gac cac ctg cct ggg gga cac act gcg cat ttg tgc agc gaa			1202
Pro Thr Asp His Leu Pro Gly Gly His Thr Ala His Leu Cys Ser Glu			
65	70	75	
aga acc ctg cga ccc ccg cct cag tct cct cag ccc gca ccc ccg ccg			1250
Arg Thr Leu Arg Pro Pro Pro Gln Ser Pro Gln Pro Ala Pro Pro Pro			
80	85	90	
cct ggt ccc gcg ctg cag tct cct ccc gct gcg ctg cgc ggg gca cgc			1298
Pro Gly Pro Ala Leu Gln Ser Pro Pro Ala Ala Leu Arg Gly Ala Arg			
95	100	105	
gcg gcg cgt gca gga acc cgg agc agc cgc gca cgg acc aca gat gcg			1346
Ala Ala Arg Ala Gly Thr Arg Ser Ser Arg Ala Arg Thr Thr Asp Ala			
110	115	120	
cgc ggc tgc cgc ctg cgc tgc cag ctg gtg ccg gtg agc gcg ctg ggc			1394
Arg Gly Cys Arg Leu Arg Ser Gln Leu Val Pro Val Ser Ala Leu Gly			
125	130	135	140
cta ggc cac agc tcc gac gag ctg ata cgt ttc cgc ttc tgc agc ggc			1442
Leu Gly His Ser Ser Asp Glu Leu Ile Arg Phe Arg Phe Cys Ser Gly			
145	150	155	
tcg tgc cgc cga gca cgc tcc cag cac gat ctg agt ctg gcc agc cta			1490
Ser Cys Arg Arg Ala Arg Ser Gln His Asp Leu Ser Leu Ala Ser Leu			
160	165	170	
ctg ggc gct ggg gcc cta cgg tgc cct ccc ggg tcc cgg ccg atc agc			1538
Leu Gly Ala Gly Ala Leu Arg Ser Pro Pro Gly Ser Arg Pro Ile Ser			
175	180	185	
cag ccc tgc tgc cgg ccc act cgc tat gag gcc gtc tcc ttc atg gac			1586
Gln Pro Cys Cys Arg Pro Thr Arg Tyr Glu Ala Val Ser Phe Met Asp			
190	195	200	
gtg aac agc acc tgg agg acc gtg gac cac ctg tcc gcc act gcc tgc			1634
Val Asn Ser Thr Trp Arg Thr Val Asp His Leu Ser Ala Thr Ala Cys			
205	210	215	220
ggc tgt ctg gcc tgaggatgat ctatctccaa gcctttgcac actagaccca			1686
Gly Cys Leu Gly			
tgtgttgccc tacctggaac agctccaccg ggctcacta accaggagcc tcaactcagc			1746
aggatatgga ggctgcagag ctccaggcccc aggcgggtga gtgacagacg tcgtcggcat			1806
gacagacaga gtgaaagatg tcggaaccac tgaccaacag tcccaagttg ttcattggatc			1866
ccagctctac agacaggaga aacctcagct aaagagaact cctctgggag aatccagaaa			1926
tggeccctctg tcttggggaa tgaattttga agagatatat atacatatat acattgtagt			1986
cgcgttgctg gaccagcctg tgctgaaacc agtcccgtgt tcaacttggtg aagccgaagc			2046

cctattttatt attttctaaat tattttattta ctttgaaaaa aaacggccaa gtcggcctcc 2106

ctttagttag gggttaatttg tgatcccggg 2136

<210> 16

<211> 224

<212> PRT

<213> Murinae gen. sp.

<400> 16

Met Glu Leu Gly Leu Ala Glu Pro Thr Ala Leu Ser His Cys Leu Arg  
1 5 10 15

Pro Arg Trp Gln Ser Ala Trp Trp Pro Thr Leu Ala Val Leu Ala Leu  
20 25 30

Leu Ser Cys Val Thr Glu Ala Ser Leu Asp Pro Met Ser Arg Ser Pro  
35 40 45

Ala Ala Arg Asp Gly Pro Ser Pro Val Leu Ala Pro Pro Thr Asp His  
50 55 60

Leu Pro Gly Gly His Thr Ala His Leu Cys Ser Glu Arg Thr Leu Arg  
65 70 75 80

Pro Pro Pro Gln Ser Pro Gln Pro Ala Pro Pro Pro Pro Gly Pro Ala  
85 90 95

Leu Gln Ser Pro Pro Ala Ala Leu Arg Gly Ala Arg Ala Ala Arg Ala  
100 105 110

Gly Thr Arg Ser Ser Arg Ala Arg Thr Thr Asp Ala Arg Gly Cys Arg  
115 120 125

Leu Arg Ser Gln Leu Val Pro Val Ser Ala Leu Gly Leu Gly His Ser  
130 135 140

Ser Asp Glu Leu Ile Arg Phe Arg Phe Cys Ser Gly Ser Cys Arg Arg  
145 150 155 160

Ala Arg Ser Gln His Asp Leu Ser Leu Ala Ser Leu Leu Gly Ala Gly  
165 170 175

Ala Leu Arg Ser Pro Pro Gly Ser Arg Pro Ile Ser Gln Pro Cys Cys  
180 185 190

Arg Pro Thr Arg Tyr Glu Ala Val Ser Phe Met Asp Val Asn Ser Thr  
195 200 205

Trp Arg Thr Val Asp His Leu Ser Ala Thr Ala Cys Gly Cys Leu Gly  
210 215 220

<210> 17

<211> 18

<212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence: PCR Primer

<400> 17

cctggccagc ctactggg

18

<210> 18  
<211> 20  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Description of Artificial Sequence: PCR Primer

<400> 18  
aaggagaccg cttcgtagcg 20

<210> 19  
<211> 17  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Description of Artificial Sequence: PCR Primer

<400> 19  
atggaacttg gacttgg 17

<210> 20  
<211> 16  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Description of Artificial Sequence: PCR Primer

<400> 20  
tccatcaccc accggc 16

<210> 21  
<211> 18  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Description of Artificial Sequence: PCR Primer

<400> 21  
ggccaccgct ccgacgag 18

<210> 22  
<211> 20  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Description of Artificial Sequence: PCR Primer

<400> 22  
ggcgggtccac ggttctccag 20

<210> 23  
<211> 29  
<212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence: PCR Primer

<400> 23

ccaagccac ctgggtgcc tctttctcc

29

<210> 24

<211> 27

<212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence: PCR Primer

<400> 24

catcaccac cggcagggc ctctcag

27

<210> 25

<211> 35

<212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence: PCR Primer

<400> 25

gagcccatgc ccggcctgat ctcagccga ggaca

35

<210> 26

<211> 34

<212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence: PCR Primer

<400> 26

ccctggctga ggccgctggc tagtgggact ctgc

34

<210> 27

<211> 31

<212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence: Hybridization Probe

<220>

<221> misc\_structure

<222> (1)

<223> wherein n represents a conjugant moiety linking to alkaline phosphatase

<400> 27

ncaggtggc cgtgggggc gccaagaccg g

31

<210> 28

<211> 16  
 <212> DNA  
 <213> Artificial Sequence

<220>  
 <223> Description of Artificial Sequence: PCR primer

<400> 28  
 ctaggagccc atgccc 16

<210> 29  
 <211> 351  
 <212> DNA  
 <213> Homo sapiens

<400> 29  
 atggctggag gaccgggac tctgtctcgt gcagcaggag cacgtggctg tctgtctcgt 60  
 tctcaactag tgcgggtgcg tgcactcggg ctgggacacc gttccgacga actagtacgt 120  
 tttcgttttt gtccaggatc ttgtcgtcgt gcacgttctc cgcgtgatct atctctagca 180  
 tctctactag gagccggagc actaagaccg ccgcccggat ctagacctgt atctcaacct 240  
 tgtttagtag ctactagata cgaagcagta tctttcatgg acgtaaactc tacatggaga 300  
 accgtagata gactatctgc aaccgcatgt ggctgtctag gatgataata g 351

<210> 30  
 <211> 414  
 <212> DNA  
 <213> Homo sapiens

<400> 30  
 atgggccatc atcatcatca tcatcatcat catcactcga gcggccatat cgacgacgac 60  
 gacaaggctg gaggaccggg atctcgtgct cgtgcagcag gagcacgtgg ctgtcgtctg 120  
 cgtttctcaac tagtgccggt gcgtgcactc ggactgggac accgttccga cgaactagta 180  
 cgttttcggt ttgtttcagg atcttctcgt cgtgcacgtt ctccgcatga tctatctcta 240  
 gcatctctac taggagccgg agcactaaga ccgcccggcg gatctagacc tgtatctcaa 300  
 ccttggtgta gacctactag atacgaagca gtatctttca tggacgtaaa ctctacatgg 360  
 agaaccgtag atagactatc tgcaaccgca tgtggctgtc taggatgata atag 414

<210> 31  
 <211> 39  
 <212> DNA  
 <213> Artificial Sequence

<220>  
 <223> Description of Artificial Sequence: PCR primer

<400> 31  
 aaggaaaaaa gcggccgcca tggaacttgg acttgagg 39

<210> 32  
 <211> 39  
 <212> DNA  
 <213> Artificial Sequence

<220>  
 <223> Description of Artificial Sequence: PCR primer

<400> 32  
 ttttttcctt ggcgccgct cagcccaggc agccgcagg 39

<210> 33

<211> 16  
 <212> DNA  
 <213> Artificial Sequence

<220>  
 <223> Description of Artificial Sequence: primer

<400> 33  
 gagcgagccc tcagcc

16

<210> 34  
 <211> 224  
 <212> PRT  
 <213> Rattus sp.

<400> 34  
 Met Glu Leu Gly Leu Gly Glu Pro Thr Ala Leu Ser His Cys Leu Arg  
           1                  5                  10                  15  
 Pro Arg Trp Gln Pro Ala Leu Trp Pro Thr Leu Ala Ala Leu Ala Leu  
                   20                  25                  30  
 Leu Ser Ser Val Thr Glu Ala Ser Leu Asp Pro Met Ser Arg Ser Pro  
                   35                  40                  45  
 Ala Ser Arg Asp Val Pro Ser Pro Val Leu Ala Pro Pro Thr Asp Tyr  
                   50                  55                  60  
 Leu Pro Gly Gly His Thr Ala His Leu Cys Ser Glu Arg Ala Leu Arg  
           65                  70                  75                  80  
 Pro Pro Pro Gln Ser Pro Gln Pro Ala Pro Pro Pro Gly Pro Ala  
                   85                  90                  95  
 Leu Gln Ser Pro Pro Ala Ala Leu Arg Gly Ala Arg Ala Ala Arg Ala  
                   100                  105                  110  
 Gly Thr Arg Ser Ser Arg Ala Arg Ala Thr Asp Ala Arg Gly Cys Arg  
                   115                  120                  125  
 Leu Arg Ser Gln Leu Val Pro Val Ser Ala Leu Gly Leu Gly His Ser  
           130                  135                  140  
 Ser Asp Glu Leu Ile Arg Phe Arg Phe Cys Ser Gly Ser Cys Arg Arg  
           145                  150                  155                  160  
 Ala Arg Ser Pro His Asp Leu Ser Leu Ala Ser Leu Leu Gly Ala Gly  
                   165                  170                  175  
 Ala Leu Arg Ser Pro Pro Gly Ser Arg Pro Ile Ser Gln Pro Cys Cys  
                   180                  185                  190  
 Arg Pro Thr Arg Tyr Glu Ala Val Ser Phe Met Asp Val Asn Ser Thr  
           195                  200                  205  
 Trp Arg Thr Val Asp His Leu Ser Ala Thr Ala Cys Gly Cys Leu Gly  
           210                  215                  220

<210> 35  
 <211> 112  
 <212> PRT  
 <213> Homo sapiens

&lt;400&gt; 35

Gly Gly Pro Gly Ser Arg Ala Arg Ala Ala Gly Ala Arg Gly Cys Arg  
 1 5 10 15  
 Leu Arg Ser Gln Leu Val Pro Val Arg Ala Leu Gly Leu Gly His Arg  
 20 25 30  
 Ser Asp Glu Leu Val Arg Phe Arg Phe Cys Ser Gly Ser Cys Arg Arg  
 35 40 45  
 Ala Arg Ser Pro His Asp Leu Ser Leu Ala Ser Leu Leu Gly Ala Gly  
 50 55 60  
 Ala Leu Arg Pro Pro Pro Gly Ser Arg Pro Val Ser Gln Pro Cys Cys  
 65 70 75 80  
 Arg Pro Thr Arg Tyr Glu Ala Val Ser Phe Met Asp Val Asn Ser Thr  
 85 90 95  
 Trp Arg Thr Val Asp Arg Leu Ser Ala Thr Ala Cys Gly Cys Leu Gly  
 100 105 110

&lt;210&gt; 36

&lt;211&gt; 111

&lt;212&gt; PRT

&lt;213&gt; Homo sapiens

&lt;400&gt; 36

Gly Pro Gly Ser Arg Ala Arg Ala Ala Gly Ala Arg Gly Cys Arg Leu  
 1 5 10 15  
 Arg Ser Gln Leu Val Pro Val Arg Ala Leu Gly Leu Gly His Arg Ser  
 20 25 30  
 Asp Glu Leu Val Arg Phe Arg Phe Cys Ser Gly Ser Cys Arg Arg Ala  
 35 40 45  
 Arg Ser Pro His Asp Leu Ser Leu Ala Ser Leu Leu Gly Ala Gly Ala  
 50 55 60  
 Leu Arg Pro Pro Pro Gly Ser Arg Pro Val Ser Gln Pro Cys Cys Arg  
 65 70 75 80  
 Pro Thr Arg Tyr Glu Ala Val Ser Phe Met Asp Val Asn Ser Thr Trp  
 85 90 95  
 Arg Thr Val Asp Arg Leu Ser Ala Thr Ala Cys Gly Cys Leu Gly  
 100 105 110

&lt;210&gt; 37

&lt;211&gt; 110

&lt;212&gt; PRT

&lt;213&gt; Homo sapiens

&lt;400&gt; 37

Pro Gly Ser Arg Ala Arg Ala Ala Gly Ala Arg Gly Cys Arg Leu Arg  
 1 5 10 15  
 Ser Gln Leu Val Pro Val Arg Ala Leu Gly Leu Gly His Arg Ser Asp  
 20 25 30  
 Glu Leu Val Arg Phe Arg Phe Cys Ser Gly Ser Cys Arg Arg Ala Arg  
 35 40 45



Ser Pro His Asp Leu Ser Leu Ala Ser Leu Leu Gly Ala Gly Ala Leu  
 50 55 60

Arg Pro Pro Pro Gly Ser Arg Pro Val Ser Gln Pro Cys Cys Arg Pro  
 65 70 75 80

Thr Arg Tyr Glu Ala Val Ser Phe Met Asp Val Asn Ser Thr Trp Arg  
 85 90 95

Thr Val Asp Arg Leu Ser Ala Thr Ala Cys Gly Cys Leu Gly  
 100 105 110

<210> 38  
 <211> 109  
 <212> PRT  
 <213> Homo sapiens

<400> 38  
 Gly Ser Arg Ala Arg Ala Ala Gly Ala Arg Gly Cys Arg Leu Arg Ser  
 1 5 10 15

Gln Leu Val Pro Val Arg Ala Leu Gly Leu Gly His Arg Ser Asp Glu  
 20 25 30

Leu Val Arg Phe Arg Phe Cys Ser Gly Ser Cys Arg Arg Ala Arg Ser  
 35 40 45

Pro His Asp Leu Ser Leu Ala Ser Leu Leu Gly Ala Gly Ala Leu Arg  
 50 55 60

Pro Pro Pro Gly Ser Arg Pro Val Ser Gln Pro Cys Cys Arg Pro Thr  
 65 70 75 80

Arg Tyr Glu Ala Val Ser Phe Met Asp Val Asn Ser Thr Trp Arg Thr  
 85 90 95

Val Asp Arg Leu Ser Ala Thr Ala Cys Gly Cys Leu Gly  
 100 105

<210> 39  
 <211> 108  
 <212> PRT  
 <213> Homo sapiens

<400> 39  
 Ser Arg Ala Arg Ala Ala Gly Ala Arg Gly Cys Arg Leu Arg Ser Gln  
 1 5 10 15

Leu Val Pro Val Arg Ala Leu Gly Leu Gly His Arg Ser Asp Glu Leu  
 20 25 30

Val Arg Phe Arg Phe Cys Ser Gly Ser Cys Arg Arg Ala Arg Ser Pro  
 35 40 45

His Asp Leu Ser Leu Ala Ser Leu Leu Gly Ala Gly Ala Leu Arg Pro  
 50 55 60

Pro Pro Gly Ser Arg Pro Val Ser Gln Pro Cys Cys Arg Pro Thr Arg  
 65 70 75 80

Tyr Glu Ala Val Ser Phe Met Asp Val Asn Ser Thr Trp Arg Thr Val  
 85 90 95

Asp Arg Leu Ser Ala Thr Ala Cys Gly Cys Leu Gly  
 100 105

<210> 40  
 <211> 107  
 <212> PRT  
 <213> Homo sapiens

<400> 40  
 Arg Ala Arg Ala Ala Gly Ala Arg Gly Cys Arg Leu Arg Ser Gln Leu  
 1 5 10 15  
 Val Pro Val Arg Ala Leu Gly Leu Gly His Arg Ser Asp Glu Leu Val  
 20 25 30  
 Arg Phe Arg Phe Cys Ser Gly Ser Cys Arg Arg Ala Arg Ser Pro His  
 35 40 45  
 Asp Leu Ser Leu Ala Ser Leu Leu Gly Ala Gly Ala Leu Arg Pro Pro  
 50 55 60  
 Pro Gly Ser Arg Pro Val Ser Gln Pro Cys Cys Arg Pro Thr Arg Tyr  
 65 70 75 80  
 Glu Ala Val Ser Phe Met Asp Val Asn Ser Thr Trp Arg Thr Val Asp  
 85 90 95  
 Arg Leu Ser Ala Thr Ala Cys Gly Cys Leu Gly  
 100 105

<210> 41  
 <211> 106  
 <212> PRT  
 <213> Homo sapiens

<400> 41  
 Ala Arg Ala Ala Gly Ala Arg Gly Cys Arg Leu Arg Ser Gln Leu Val  
 1 5 10 15  
 Pro Val Arg Ala Leu Gly Leu Gly His Arg Ser Asp Glu Leu Val Arg  
 20 25 30  
 Phe Arg Phe Cys Ser Gly Ser Cys Arg Arg Ala Arg Ser Pro His Asp  
 35 40 45  
 Leu Ser Leu Ala Ser Leu Leu Gly Ala Gly Ala Leu Arg Pro Pro Pro  
 50 55 60  
 Gly Ser Arg Pro Val Ser Gln Pro Cys Cys Arg Pro Thr Arg Tyr Glu  
 65 70 75 80  
 Ala Val Ser Phe Met Asp Val Asn Ser Thr Trp Arg Thr Val Asp Arg  
 85 90 95  
 Leu Ser Ala Thr Ala Cys Gly Cys Leu Gly  
 100 105

<210> 42  
 <211> 105  
 <212> PRT  
 <213> Homo sapiens

&lt;400&gt; 42

Arg Ala Ala Gly Ala Arg Gly Cys Arg Leu Arg Ser Gln Leu Val Pro  
 1 5 10 15  
 Val Arg Ala Leu Gly Leu Gly His Arg Ser Asp Glu Leu Val Arg Phe  
 20 25 30  
 Arg Phe Cys Ser Gly Ser Cys Arg Arg Ala Arg Ser Pro His Asp Leu  
 35 40 45  
 Ser Leu Ala Ser Leu Leu Gly Ala Gly Ala Leu Arg Pro Pro Pro Gly  
 50 55 60  
 Ser Arg Pro Val Ser Gln Pro Cys Cys Arg Pro Thr Arg Tyr Glu Ala  
 65 70 75 80  
 Val Ser Phe Met Asp Val Asn Ser Thr Trp Arg Thr Val Asp Arg Leu  
 85 90 95  
 Ser Ala Thr Ala Cys Gly Cys Leu Gly  
 100 105

&lt;210&gt; 43

&lt;211&gt; 104

&lt;212&gt; PRT

&lt;213&gt; Homo sapiens

&lt;400&gt; 43

Ala Ala Gly Ala Arg Gly Cys Arg Leu Arg Ser Gln Leu Val Pro Val  
 1 5 10 15  
 Arg Ala Leu Gly Leu Gly His Arg Ser Asp Glu Leu Val Arg Phe Arg  
 20 25 30  
 Phe Cys Ser Gly Ser Cys Arg Arg Ala Arg Ser Pro His Asp Leu Ser  
 35 40 45  
 Leu Ala Ser Leu Leu Gly Ala Gly Ala Leu Arg Pro Pro Pro Gly Ser  
 50 55 60  
 Arg Pro Val Ser Gln Pro Cys Cys Arg Pro Thr Arg Tyr Glu Ala Val  
 65 70 75 80  
 Ser Phe Met Asp Val Asn Ser Thr Trp Arg Thr Val Asp Arg Leu Ser  
 85 90 95  
 Ala Thr Ala Cys Gly Cys Leu Gly  
 100

&lt;210&gt; 44

&lt;211&gt; 103

&lt;212&gt; PRT

&lt;213&gt; Homo sapiens

&lt;400&gt; 44

Ala Gly Ala Arg Gly Cys Arg Leu Arg Ser Gln Leu Val Pro Val Arg  
 1 5 10 15  
 Ala Leu Gly Leu Gly His Arg Ser Asp Glu Leu Val Arg Phe Arg Phe  
 20 25 30  
 Cys Ser Gly Ser Cys Arg Arg Ala Arg Ser Pro His Asp Leu Ser Leu

35 40 45  
 Ala Ser Leu Leu Gly Ala Gly Ala Leu Arg Pro Pro Pro Gly Ser Arg  
 50 55 60  
 Pro Val Ser Gln Pro Cys Cys Arg Pro Thr Arg Tyr Glu Ala Val Ser  
 65 70 75 80  
 Phe Met Asp Val Asn Ser Thr Trp Arg Thr Val Asp Arg Leu Ser Ala  
 85 90 95  
 Thr Ala Cys Gly Cys Leu Gly  
 100

<210> 45  
 <211> 102  
 <212> PRT  
 <213> Homo sapiens

<400> 45  
 Gly Ala Arg Gly Cys Arg Leu Arg Ser Gln Leu Val Pro Val Arg Ala  
 1 5 10 15  
 Leu Gly Leu Gly His Arg Ser Asp Glu Leu Val Arg Phe Arg Phe Cys  
 20 25 30  
 Ser Gly Ser Cys Arg Arg Ala Arg Ser Pro His Asp Leu Ser Leu Ala  
 35 40 45  
 Ser Leu Leu Gly Ala Gly Ala Leu Arg Pro Pro Pro Gly Ser Arg Pro  
 50 55 60  
 Val Ser Gln Pro Cys Cys Arg Pro Thr Arg Tyr Glu Ala Val Ser Phe  
 65 70 75 80  
 Met Asp Val Asn Ser Thr Trp Arg Thr Val Asp Arg Leu Ser Ala Thr  
 85 90 95  
 Ala Cys Gly Cys Leu Gly  
 100

<210> 46  
 <211> 101  
 <212> PRT  
 <213> Homo sapiens

<400> 46  
 Ala Arg Gly Cys Arg Leu Arg Ser Gln Leu Val Pro Val Arg Ala Leu  
 1 5 10 15  
 Gly Leu Gly His Arg Ser Asp Glu Leu Val Arg Phe Arg Phe Cys Ser  
 20 25 30  
 Gly Ser Cys Arg Arg Ala Arg Ser Pro His Asp Leu Ser Leu Ala Ser  
 35 40 45  
 Leu Leu Gly Ala Gly Ala Leu Arg Pro Pro Pro Gly Ser Arg Pro Val  
 50 55 60  
 Ser Gln Pro Cys Cys Arg Pro Thr Arg Tyr Glu Ala Val Ser Phe Met  
 65 70 75 80  
 Asp Val Asn Ser Thr Trp Arg Thr Val Asp Arg Leu Ser Ala Thr Ala

85 90 95

Cys Gly Cys Leu Gly  
100

<210> 47  
<211> 100  
<212> PRT  
<213> Homo sapiens

<400> 47  
Arg Gly Cys Arg Leu Arg Ser Gln Leu Val Pro Val Arg Ala Leu Gly  
1 5 10 15  
Leu Gly His Arg Ser Asp Glu Leu Val Arg Phe Arg Phe Cys Ser Gly  
20 25 30  
Ser Cys Arg Arg Ala Arg Ser Pro His Asp Leu Ser Leu Ala Ser Leu  
35 40 45  
Leu Gly Ala Gly Ala Leu Arg Pro Pro Pro Gly Ser Arg Pro Val Ser  
50 55 60  
Gln Pro Cys Cys Arg Pro Thr Arg Tyr Glu Ala Val Ser Phe Met Asp  
65 70 75 80  
Val Asn Ser Thr Trp Arg Thr Val Asp Arg Leu Ser Ala Thr Ala Cys  
85 90 95  
Gly Cys Leu Gly  
100

<210> 48  
<211> 99  
<212> PRT  
<213> Homo sapiens

<400> 48  
Gly Cys Arg Leu Arg Ser Gln Leu Val Pro Val Arg Ala Leu Gly Leu  
1 5 10 15  
Gly His Arg Ser Asp Glu Leu Val Arg Phe Arg Phe Cys Ser Gly Ser  
20 25 30  
Cys Arg Arg Ala Arg Ser Pro His Asp Leu Ser Leu Ala Ser Leu Leu  
35 40 45  
Gly Ala Gly Ala Leu Arg Pro Pro Pro Gly Ser Arg Pro Val Ser Gln  
50 55 60  
Pro Cys Cys Arg Pro Thr Arg Tyr Glu Ala Val Ser Phe Met Asp Val  
65 70 75 80  
Asn Ser Thr Trp Arg Thr Val Asp Arg Leu Ser Ala Thr Ala Cys Gly  
85 90 95  
Cys Leu Gly

<210> 49  
<211> 197  
<212> PRT

&lt;213&gt; Homo sapiens

&lt;400&gt; 49

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Met Gln Arg Trp Lys Ala Ala Ala Leu Ala Ser Val Leu Cys Ser Ser
 1           5           10           15

Val Leu Ser Ile Trp Met Cys Arg Glu Gly Leu Leu Leu Ser His Arg
 20           25           30

Leu Gly Pro Ala Leu Val Pro Leu His Arg Leu Pro Arg Thr Leu Asp
 35           40           45

Ala Arg Ile Ala Arg Leu Ala Gln Tyr Arg Ala Leu Leu Gln Gly Ala
 50           55           60

Pro Asp Ala Met Glu Leu Arg Glu Leu Thr Pro Trp Ala Gly Arg Pro
 65           70           75           80

Pro Gly Pro Arg Arg Arg Ala Gly Pro Arg Arg Arg Arg Ala Arg Ala
 85           90           95

Arg Leu Gly Ala Arg Pro Cys Gly Leu Arg Glu Leu Glu Val Arg Val
100           105           110

Ser Glu Leu Gly Leu Gly Tyr Ala Ser Asp Glu Thr Val Leu Phe Arg
115           120           125

Tyr Cys Ala Gly Ala Cys Glu Ala Ala Ala Arg Val Tyr Asp Leu Gly
130           135           140

Leu Arg Arg Leu Arg Gln Arg Arg Arg Leu Arg Arg Glu Arg Val Arg
145           150           155           160

Ala Gln Pro Cys Cys Arg Pro Thr Ala Tyr Glu Asp Glu Val Ser Phe
165           170           175

Leu Asp Ala His Ser Arg Tyr His Thr Val His Glu Leu Ser Ala Arg
180           185           190

Glu Cys Ala Cys Val
195

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&lt;210&gt; 50

&lt;211&gt; 156

&lt;212&gt; PRT

&lt;213&gt; Homo sapiens

&lt;400&gt; 50

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Met Ala Val Gly Lys Phe Leu Leu Gly Ser Leu Leu Leu Leu Ser Leu
 1           5           10           15

Gln Leu Gly Gln Gly Trp Gly Pro Asp Ala Arg Gly Val Pro Val Ala
 20           25           30

Asp Gly Glu Phe Ser Ser Glu Gln Val Ala Lys Ala Gly Gly Thr Trp
 35           40           45

Leu Gly Thr His Arg Pro Leu Ala Arg Leu Arg Arg Ala Leu Ser Gly
 50           55           60

Pro Cys Gln Leu Trp Ser Leu Thr Leu Ser Val Ala Glu Leu Gly Leu
 65           70           75           80

Gly Tyr Ala Ser Glu Glu Lys Val Ile Phe Arg Tyr Cys Ala Gly Ser

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85 90 95  
 Cys Pro Arg Gly Ala Arg Thr Gln His Gly Leu Ala Leu Ala Arg Leu  
 100 105 110  
 Gln Gly Gln Gly Arg Ala His Gly Gly Pro Cys Cys Arg Pro Thr Arg  
 115 120 125  
 Tyr Thr Asp Val Ala Phe Leu Asp Asp Arg His Arg Trp Gln Arg Leu  
 130 135 140  
 Pro Gln Leu Ser Ala Ala Ala Cys Gly Cys Gly Gly  
 145 150 155

<210> 51  
 <211> 211  
 <212> PRT  
 <213> Homo sapiens

<400> 51  
 Met Lys Leu Trp Asp Val Val Ala Val Cys Leu Val Leu Leu His Thr  
 1 5 10 15  
 Ala Ser Ala Phe Pro Leu Pro Ala Gly Lys Arg Pro Pro Glu Ala Pro  
 20 25 30  
 Ala Glu Asp Arg Ser Leu Gly Arg Arg Arg Ala Pro Phe Ala Leu Ser  
 35 40 45  
 Ser Asp Ser Asn Met Pro Glu Asp Tyr Pro Asp Gln Phe Asp Asp Val  
 50 55 60  
 Met Asp Phe Ile Gln Ala Thr Ile Lys Arg Leu Lys Arg Ser Pro Asp  
 65 70 75 80  
 Lys Gln Met Ala Val Leu Pro Arg Arg Glu Arg Asn Arg Gln Ala Ala  
 85 90 95  
 Ala Ala Asn Pro Glu Asn Ser Arg Gly Lys Gly Arg Arg Gly Gln Arg  
 100 105 110  
 Gly Lys Asn Arg Gly Cys Val Leu Thr Ala Ile His Leu Asn Val Thr  
 115 120 125  
 Asp Leu Gly Leu Gly Tyr Glu Thr Lys Glu Glu Leu Ile Phe Arg Tyr  
 130 135 140  
 Cys Ser Gly Ser Cys Asp Ala Ala Glu Thr Thr Tyr Asp Lys Ile Leu  
 145 150 155 160  
 Lys Asn Leu Ser Arg Asn Arg Arg Leu Val Ser Asp Lys Val Gly Gln  
 165 170 175  
 Ala Cys Cys Arg Pro Ile Ala Phe Asp Asp Asp Leu Ser Phe Leu Asp  
 180 185 190  
 Asp Asn Leu Val Tyr His Ile Leu Arg Lys His Ser Ala Lys Arg Cys  
 195 200 205  
 Gly Cys Ile  
 210

<210> 52

<211> 365  
 <212> DNA  
 <213> Artificial Sequence

<220>  
 <223> Description of Artificial Sequence:synthetic gene  
 for Neublastin

<400> 52  
 taccatggct ggaggaccgg gatctcgtgc tcgtgcagca ggagcacgtg gctgtcgtct 60  
 gcgtttctcaa ctagtgccgg tgcgtgcact cggactggga caccgttccg acgaactagt 120  
 acgttttcgt ttttgttcag gatcttgtcg tcgtgcacgt tctccgcatg atctatctct 180  
 agcatctcta ctaggagccg gagcactaag accgccgccg ggatctagac ctgtatctca 240  
 accttgttgt agacctacta gatacgaagc agtatctttc atggacgtaa actctacatg 300  
 gagaaccgta gatagactat ctgcaaccgc atgtggctgt ctaggatgat aatagggatc 360  
 cggt 365

<210> 53  
 <211> 365  
 <212> DNA  
 <213> Artificial Sequence

<220>  
 <223> Description of Artificial Sequence:synthetic gene  
 for Neublastin

<400> 53  
 atggtaccga cctcctggcc ctagagcacg agcacgtcgt cctcgtgcac cgacagcaga 60  
 cgcaagagtt gatcacggcc acgcacgtga gcctgacct gtggcaaggc tgcttgatca 120  
 tgcaaaagca aaaacaagtc ctagaacagc agcacgtgca agaggcgtag tagatagaga 180  
 tcgtagagat gatcctcgcc ctctgtgattc tggcggcggc cctagatctg gacatagagt 240  
 tggaacaaca tctggatgat ctatgcttcg tcatagaaag tacctgcatt tgagatgtac 300  
 ctcttggcat ctatctgata gacgttggcg tacaccgaca gatcctacta ttatccctag 360  
 gccga 365

<210> 54  
 <211> 114  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Description of Artificial Sequence:synthetic  
 Neublastin

<400> 54  
 Met Ala Gly Gly Pro Gly Ser Arg Ala Arg Ala Ala Gly Ala Arg Gly  
 1 5 10 15  
 Cys Arg Leu Arg Ser Gln Leu Val Pro Val Arg Ala Leu Gly Leu Gly  
 20 25 30  
 His Arg Ser Asp Glu Leu Val Arg Phe Arg Phe Cys Ser Gly Ser Cys  
 35 40 45  
 Arg Arg Ala Arg Ser Pro His Asp Leu Ser Leu Ala Ser Leu Leu Gly  
 50 55 60  
 Ala Gly Ala Leu Arg Pro Pro Pro Gly Ser Arg Pro Val Ser Gln Pro  
 65 70 75 80  
 Cys Cys Arg Pro Thr Arg Tyr Glu Ala Val Ser Phe Met Asp Val Asn  
 85 90 95



Ser Thr Trp Arg Thr Val Asp Arg Leu Ser Ala Thr Ala Cys Gly Cys  
 100 105 110

Leu Gly

<210> 55  
 <211> 442  
 <212> DNA  
 <213> Artificial Sequence

<220>  
 <223> Description of Artificial Sequence:synthetic gene  
 for HisNeublastin

<400> 55  
 taccatgggc catcatcatc atcatcatca tcatcatcac tcgagcggcc atatcgacga 60  
 cgacgacaag gctggaggac cgggatctcg tgctcgtgca gcaggagcac gtggctgtcg 120  
 tctgcgttct caactagtgc cgggtcgtgc actcggactg ggacaccggt cgcacgaact 180  
 agtacgtttt cgtttttgtt caggatcttg tcgtcgtgca cgttctccgc atgatctatc 240  
 tctagcatct ctactaggag ccggagcact aagaccgccg ccgggatcta gacctgtatc 300  
 tcaaccttgt ttagaccta ctagatacga agcagtatct ttcatggacg taaactctac 360  
 atggagaacc gtagatagac tatctgcaac cgcagtgggc tgtctaggat gataataggg 420  
 atccggctgc taacaaagcc cg 442

<210> 56  
 <211> 442  
 <212> DNA  
 <213> Artificial Sequence

<220>  
 <223> Description of Artificial Sequence:synthetic gene  
 for HisNeublastin

<400> 56  
 atggtaccgc gtagtagtag tagtagtagt agtagtagtg agctcgccgg tatagctgct 60  
 gctgctgttc cgacctctcg gccctagagc acgagcacgt cgtcctcgtg caccgacagc 120  
 agacgcaaga gttgatcacg gccacgcacg tgagcctgac cctgtggcaa ggctgcttga 180  
 tcatgcaaaa gcaaaaacaa gtccatagaac agcagcacgt gcaagaggcg tactagatag 240  
 agatcgtaga gatgatcctc ggccctcgtg ttctggcggc ggccctagat ctggacatag 300  
 agttggaaca acatctggat gatctatgct tcgtcataga aagtacctgc atttgagatg 360  
 tacctcttgg catctatctg atagacgttg gcgtacaccg acagatccta ctattatccc 420  
 taggcgacg attgttcgg gc 442

<210> 57  
 <211> 135  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Description of Artificial Sequence:synthetic  
 HisNeublastin

<400> 57  
 Met Gly His His His His His His His His His Ser Ser Gly His  
 1 5 10 15  
 Ile Asp Asp Asp Asp Lys Ala Gly Gly Pro Gly Ser Arg Ala Arg Ala  
 20 25 30  
 Ala Gly Ala Arg Gly Cys Arg Leu Arg Ser Gln Leu Val Pro Val Arg  
 35 40 45

Ala Leu Gly Leu Gly His Arg Ser Asp Glu Leu Val Arg Phe Arg Phe  
50 55 60

Cys Ser Gly Ser Cys Arg Arg Ala Arg Ser Pro His Asp Leu Ser Leu  
65 70 75 80

Ala Ser Leu Leu Gly Ala Gly Ala Leu Arg Pro Pro Pro Gly Ser Arg  
85 90 95

Pro Val Ser Gln Pro Cys Cys Arg Pro Thr Arg Tyr Glu Ala Val Ser  
100 105 110

Phe Met Asp Val Asn Ser Thr Trp Arg Thr Val Asp Arg Leu Ser Ala  
115 120 125

Thr Ala Cys Gly Cys Leu Gly  
130 135

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